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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:02:27 ; Search time 51.308 Seconds
(without alignments)
699.375 Million cell updates/sec

Title: US-10-661-784-3
Perfect score: 697
Sequence: 1 GSKGFVQPPTKICVQCPD.....VPEKKIYFTVTVNHWECEP 127

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A Geneseq 29Jan04:*
1: Geneseqp1980s:*
2: Geneseqp1990s:*
3: Geneseqp2000s:*
4: Geneseqp2001s:*
5: Geneseqp2002s:*
6: Geneseqp2003as:*
7: Geneseqp2003bs:*
8: Geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
1	618	90.0	123	3	AA95426 Human hig
2	618	90.0	304	6	ABP70801 Human ext
3	618	90.0	322	6	ABP70799 Human ext
4	618	90.0	329	6	ABU92044 Human pro
5	618	90.0	358	6	ABP70800 Human ext
6	618	90.0	390	6	ABU99149 Novel hum
7	618	90.0	398	6	ABU99143 Novel hum
8	618	90.0	427	8	ADE76864 Human pro
9	618	90.0	615	6	ABU99144 Novel hum
10	618	90.0	626	5	ABP78707 Human hig
11	618	90.0	644	4	ABG21101 Novel hum
12	618	90.0	644	5	ABP78710 Human hig
13	618	90.0	644	6	ABU99150 Novel hum
14	618	90.0	644	6	ABU99145 Novel hum
15	596	85.3	122	3	ABP37447 Human kin
16	585	85.2	435	4	ABG21105 Novel hum
17	556.5	81.0	117	2	AA933350 Domaine 3
18	440	64.0	436	1	AA940257 Bradykini
19	413	60.1	434	1	AA940633 Bradykini
20	411	59.8	357	6	ABR41302 Human DIT
21	388	56.5	235	5	ABG60077 Human DIT
22	320.5	46.7	248	4	ABG21102 Novel hum
23	316	46.0	369	4	ABG21099 Novel hum
24	190	27.7	305	4	ABG21100 Novel hum
25	171.5	25.0	167	2	AA9898907 Mouse IMC

26	166	24.2	32	3	AA95418 Anti-angi
27	163.5	23.8	126	3	AB37445 Human cys
28	163.5	23.8	145	2	AA32223 Mature hu
29	163.5	23.8	145	2	AA31302 Human cys
30	163.5	23.8	145	2	AA925708 Human cys
31	163.5	23.8	145	4	AAE02410 Human cys
32	163.5	23.8	145	4	AAE04439 Human cys
33	163.5	23.8	145	7	ADD14098 Human src
34	163.5	23.8	167	2	AA92287 Secreted
35	163	23.7	167	2	ADA45154 Human pol
36	163	23.7	178	2	AAW69734 Human cys
37	161	23.4	32	3	AA95408 Anti-angi
38	157	22.9	122	3	AA37446 Human kin
39	154.5	22.5	167	2	AAW98910 Mouse IMC
40	153	22.3	27	3	AA95425 Anti-angi
41	144.5	21.0	121	3	AA981200 Human mut
42	144.5	21.0	128	3	AA981189 Human mut
43	143.5	20.9	121	3	AA981198 Human mut
44	143.5	20.9	128	3	AA981187 Human mut
45	142.5	20.7	118	3	AA981218 Bovine mu

ALIGNMENTS

RESULT 1
AA95426
ID AA95426 standard; peptide; 123 AA.
XX AC AA95426;
DT 25-SEP-2000 (first entry)
XX DE Human high mol.wt. kininogen domain 3.
XX KW Anti-angiogenic; angiogenesis; inhibitor; kininogen; homologue;
KW endothelial cell proliferation; apoptosis; cancer; ocular disorder;
KW rheumatoid arthritis; cytostatic; antiarthritic; antitumour; therapy;
KW human; D3 peptide.
XX OS Homo sapiens.
XX PN WO200035407-A2.
XX PD 22-JUN-2000.
XX PF 02-DEC-1999; 99WO-US028465.
XX PR 16-DEC-1998; 98US-0112427P.
XX PA (UTEM) UNIV TEMPLE.
XX PA (MCCR/) MCCRAE R K.
XX PI McCrae RK;
XX DR WPI; 2000-442247/38.
XX PT Composition for inhibiting angiogenesis and endothelial cell
PT proliferation, inducing endothelial cell apoptosis and creating cancer, 3
PT rheumatoid arthritis, and ocular disorders comprises a kininogen domain 3
PT analog.
XX PS Disclosure; Page 4; 44pp; English.
XX CC The present sequence is that of domain 3 of human high mol.wt. kininogen
CC (HK) . The invention provides peptides (see AA95405-24) that are
CC analogues of certain sites in the HK domain 3, specifically Asn275-
CC Lys282, Cys246-Cys249, Leu331-Tyr338 and Tyr299-Ser314. The peptides, in
CC which native Cys residues may be replaced by Ala residues, inhibit
CC endothelial cell proliferation and may also induce endothelial cell
CC apoptosis. Compositions including the peptides are used in claimed
CC methods for inhibiting angiogenesis, inhibiting endothelial cell
CC proliferation, and inducing endothelial cell apoptosis. Cancer,

CC rheumatoid arthritis, and ocular disorders characterized by undesired
 CC vascularization of the retina are treated
 XX
 SQ Sequence 123 AA;
 Query Match 90.0%; Score 618; DB 3; Length 123;
 Best Local Similarity 100.0%; Pred. No. 5.5e-63;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPNTPSPLEETLTHITIKLNAENNATFYFKIDNVKKARVQV 62
 DB 1 GKDFVQPTKICVGCPRDIPNTPSPLEETLTHITIKLNAENNATFYFKIDNVKKARVQV 60
 QY 63 AGKKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEYVYVVPWEKKIYPTV 118
 DB 61 AGKKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEYVYVVPWEKKIYPTV 116
 RESULT 2
 ABP70801
 ID ABP70801 standard; protein; 304 AA.
 XX
 AC ABP70801;
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-28.
 XX
 KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.
 XX
 PA (INCY-) INCYTE GENOMICS INC.
 XX
 PI Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebajadian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Ramkumar J;
 XX
 DR WPI; 2003-278643/27.
 XX
 DR N-PSDB; ACC42388.
 XX
 PT New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 207; 224pp; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP7074-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
 CC for diagnosing or treating a disease or condition associated with
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer

XX
 SQ Sequence 304 AA;
 Query Match 90.0%; Score 618; DB 6; Length 304;
 Best Local Similarity 100.0%; Pred. No. 1.9e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPNTPSPLEETLTHITIKLNAENNATFYFKIDNVKKARVQV 62
 DB 130 GKDFVQPTKICVGCPRDIPNTPSPLEETLTHITIKLNAENNATFYFKIDNVKKARVQV 189
 QY 63 AGKKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEYVYVVPWEKKIYPTV 118
 DB 190 AGKKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEYVYVVPWEKKIYPTV 245
 RESULT 3
 ABP70799
 ID ABP70799 standard; protein; 322 AA.
 XX
 AC ABP70799;
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-26.
 XX
 KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.
 XX
 PA (INCY-) INCYTE GENOMICS INC.
 XX
 PI Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebajadian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Ramkumar J;
 XX
 DR WPI; 2003-278643/27.
 XX
 DR N-PSDB; ACC42388.
 XX
 PT New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 205-206; 224pp; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP7074-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
 CC for diagnosing or treating a disease or condition associated with
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer

Sequence 322 AA;

CC	polynucleotides. The diseases or conditions associated with decreased expression or overexpression of PMM are cell proliferation disorders (e.g. cancer, atherosclerosis), neurological disorders (e.g. epilepsy, Huntington's disease, stroke), immune/inflammatory disorders (e.g. AIDS, allergies), developmental disorders (e.g. hypothyroidism, Cushing's syndrome), gastrointestinal or epithelial disorders, and infections. The PMM polypeptides or their fragments are useful in screening compounds for effectiveness as agonists or antagonists of the polypeptides, or in altering the expression of the target polynucleotide and compounds that specifically bind to, or modulate the activity of the polypeptide.
CC	ABU92021-ABU92060 represent the human PMM polypeptides of the invention
XX	
SQ	Sequence 329 AA;
	Query Match 90.0%; Score 618; DB 6; Length 329;
	Best Local Similarity 100.0%; Pred. No. 2.1e-62;
	Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	3 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITIKLNAENNAFFYKIDNVKKARQVQV 62
DB	155 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITIKLNAENNAFFYKIDNVKKARQVQV 214
QY	63 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNNAEVYVVPWEKKIYPTV 118
DB	215 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNNAEVYVVPWEKKIYPTV 270
RESULT 5	
ABP70800	
ID	ABP70800 standard; protein; 358 AA.
XX	ABP70800;
XX	26-AUG-2003 (first entry)
DT	Human extracellular messenger, EXMES-27.
DE	
XX	Human, extracellular messenger; EXMES; cytostatic; antidiabetic;
KW	Immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
KW	endocrine disorder; cancer.
XX	
OS	Homo sapiens.
XX	
XX	WO2003018612-A2.
PN	
PD	06-MAR-2003.
XX	
PF	22-AUG-2002; 2002WO-US027213.
XX	
PR	24-AUG-2001; 2001US-0314811P.
PR	14-DEC-2001; 2001US-0340584P.
PR	18-JAN-2002; 2002US-0350595P.
PR	11-MAR-2002; 2002US-0363432P.
PR	15-MAR-2002; 2002US-0364607P.
PR	05-APR-2002; 2002US-0370761P.
PR	24-JUN-2002; 2002US-0391378P.
XX	
PA	(INCY-) INCYTE GENOMICS INC.
XX	
PI	Duggan BM, Lee S, Baughn MR, Hafalia AJA, Walia NK, Elliott VS;
PI	Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang Y, Burford N;
PI	Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebartadian Y;
PI	Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
PI	Ramkumar J;
XX	
DR	WPI; 2003-278643/27.
DR	N-PSDB; ACC42387.
XX	
XX	New human extracellular messenger (EXMES) polypeptide, useful for
PT	preparing a composition for treating a disease associated with decreased
PT	expression or overexpression of functional EXMES e.g. autoimmune
PT	disorders or cancer.
XX	

QY	3 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITIKLNAENNAFFYKIDNVKKARQVQV 62
DB	148 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITIKLNAENNAFFYKIDNVKKARQVQV 207
QY	63 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNNAEVYVVPWEKKIYPTV 118
DB	208 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNNAEVYVVPWEKKIYPTV 263
RESULT 4	
ABU92044	
ID	ABU92044 standard; protein; 329 AA.
XX	ABU92044;
XX	15-JUL-2003 (first entry)
DT	
DE	Human protein modification and maintenance molecule-24 (PMM-24).
XX	
XX	Human; protein modification and maintenance molecule; PMM; cancer;
KW	cell proliferation disorder; atherosclerosis; neurological disorder;
KW	epilepsy; Huntington's disease; stroke; immune disorder; allergy;
KW	Inflammatory disorder; AIDS; developmental disorder; hypothyroidism;
KW	Cushing's syndrome; gastrointestinal disorder; epithelial disorder;
KW	infection; cytostatic; antiatherosclerotic; anticonvulsant; neurotropic;
KW	neuroprotective; cerebroprotective; anti-HIV; anti-allergic; vulnary;
KW	antiinflammatory; thyromimetic.
XX	
OS	Homo sapiens.
XX	
XX	WO2003031939-A2.
PN	
PD	17-APR-2003.
XX	
PF	11-OCT-2002; 2002WO-US032850.
XX	
PR	12-OCT-2001; 2001US-0329689P.
PR	25-OCT-2001; 2001US-0335703P.
PR	09-NOV-2001; 2001US-0348887P.
PR	28-NOV-2001; 2001US-0334145P.
PR	06-DEC-2001; 2001US-0337451P.
PR	14-DEC-2001; 2001US-0340584P.
XX	
PA	(INCY-) INCYTE GENOMICS INC.
XX	
PI	Ramkumar J, Gorvay AE, Baughn MR, Emerling BM, Yang J, Lee SY;
PI	Tran UK, Becha SD, Duggan BM, Lee EA, Griffin JA, Li JX;
PI	Sprague WM, Hafalia AJA, Chawla NK, Lehr-Mason PM, Kable AE, Yue H;
PI	Marquis JP, Yao MG, Richardson TW, Tang TY, Jin P, Chien D;
PI	Bhatia U, Burrill JD, Lee S, Blake JU, Ho A, Zheng W;
XX	
DR	WPI; 2003-430274/40.
DR	N-PSDB; ACA92439.
XX	
XX	New human protein modification and maintenance molecules (PMM), useful
PT	for diagnosing, treating and preventing diseases or conditions associated
PT	with the aberrant PMM expression e.g. cancer, atherosclerosis, or
PT	infections.
XX	
PS	Claim 1; Page 264-265; 31pp; English.
XX	
CC	The present invention relates to the isolation of human protein
CC	modification and maintenance molecules (PMM), and the polynucleotide
CC	sequences encoding them. A total of 40 PMM polypeptides (designated PMM
CC	-1 to PMM-40) are disclosed. The sequences of the invention are useful
CC	for diagnosing a condition or disease associated with the expression of
CC	PMM in a subject, preparing a polyclonal or monoclonal antibody, and
CC	generating an expression profile of a sample containing the

PS Claim 1; Page 206; 224pp; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP70774-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
 CC for diagnosing or treating a disease or condition associated with
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer
 XX
 SQ Sequence 358 AA;
 Query Match 90.0%; Score 618; DB 6; Length 358;
 Best Local Similarity 100.0%; Pred. No. 2.4e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQV 62
 Db 184 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQV 243
 QY 63 AGKYFIDFVARETTCSKESNEELTESCETKKGSLDCNAEYVVPWEKIYPTV 118
 Db 244 AGKYFIDFVARETTCSKESNEELTESCETKKGSLDCNAEYVVPWEKIYPTV 299
 RESULT 6
 ABU99149
 ID ABU99149 standard; protein; 390 AA.
 AC ABU99149;
 XX
 DT 01-AUG-2003 (first entry)
 DE Novel human GPCR related protein NOV12g.
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiant; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOV-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 OS Homo sapiens.
 XX
 PN WO200299116-A2.
 XX
 PD 12-DEC-2002.
 XX
 PF 04-JUN-2002; 2002WO-US017428.
 PR 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295661P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 06-JUN-2001; 2001US-0296418P.
 PR 14-JUN-2001; 2001US-0298285P.
 PR 15-JUN-2001; 2001US-0298556P.
 PR 21-JUN-2001; 2001US-029949P.
 PR 26-JUN-2001; 2001US-0300883P.
 PR 28-JUN-2001; 2001US-0301550P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 27-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-0322293P.
 PR 17-SEP-2001; 2001US-0322706P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR
 XX
 PR 03-JUN-2002; 2002US-00363676.
 PA (CURA-) CURAGEN CORP.
 XX Anderson DW, Baumgartner JC, Boldog FL, Casman SU, Edinger SR;
 XX Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalte T, Kekuda R, Li L;
 XX Macdougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
 XX Pena CE, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CAM;
 XX Vos EZ, Zernhusen BD;
 XX WPI; 2003-140627/13.
 DR N-PSDB; ACD03653.
 XX
 XX New NOVX polypeptides and nucleic acids, useful for preventing or
 XX treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 XX atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 XX pharmacogenomics.
 XX
 PS Claim 1; Page 147; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 SQ Sequence 390 AA;
 Query Match 90.0%; Score 618; DB 6; Length 390;
 Best Local Similarity 100.0%; Pred. No. 2.7e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQV 62
 Db 216 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQV 275
 QY 63 AGKYFIDFVARETTCSKESNEELTESCETKKGSLDCNAEYVVPWEKIYPTV 118
 Db 276 AGKYFIDFVARETTCSKESNEELTESCETKKGSLDCNAEYVVPWEKIYPTV 331
 RESULT 7
 ABU99143
 ID ABU99143 standard; protein; 398 AA.
 XX
 AC ABU99143;
 XX
 DT 01-AUG-2003 (first entry)
 DE Novel human GPCR related protein NOV12a.
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiant; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOV-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW

KW diabetes; immune disorder; AIDS; obesity; asthma;
KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
KW infection; multiple sclerosis; cancer-associated cachexia;
KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
XX Homo sapiens.
XX
XX WO200299116-A2.
XX
XX 12-DEC-2002.
XX
XX 04-JUN-2002; 2002WO-US017428.
XX
XX 04-JUN-2001; 2001US-0295607P.
XX 04-JUN-2001; 2001US-0295661P.
XX 06-JUN-2001; 2001US-0296404P.
XX 06-JUN-2001; 2001US-0296418P.
XX 14-JUN-2001; 2001US-0298285P.
XX 15-JUN-2001; 2001US-0298556P.
XX 21-JUN-2001; 2001US-0299949P.
XX 26-JUN-2001; 2001US-0300883P.
XX 28-JUN-2001; 2001US-0301550P.
XX 13-AUG-2001; 2001US-0311972P.
XX 27-AUG-2001; 2001US-0315071P.
XX 29-AUG-2001; 2001US-0315660P.
XX 14-SEP-2001; 2001US-0322293P.
XX 17-SEP-2001; 2001US-0322706P.
XX 18-DEC-2001; 2001US-0341189P.
XX 28-FEB-2002; 2002US-0361186P.
XX 12-MAR-2002; 2002US-0363673P.
XX 12-MAR-2002; 2002US-0363676P.
XX 03-JUN-2002; 2002US-00363676.
XX (CURA-) CURAGEN CORP.
XX
XX Anderson DW, Baumgartner JC, Boldog FB, Casman SJ, Edinger SR;
XX Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalt T, Kekuda R, Li L;
XX MacDougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
XX Pena CEA, Rastelli L, Shimkets RA, Stone DJ, Spytek KA, Vernet CAM;
XX Voss EZ, Zerhusen BD;
XX
XX WPI; 2003-140627/13.
XX N-PSDB; ACD03647.
XX

New NOVX polypeptides and nucleic acids, useful for preventing or treating NOVX-associated disorders, e.g. cancer, cardiomyopathy, atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or pharmacogenomics.

Claim 1; Page 143; 332pp; English.

The invention describes an isolated polypeptide (I) comprising any of 27 118-961 residue amino acid sequences, given in the specification, a mature form of them, a sequence that is at least 95 % identical to them, or a sequence having one or more conservative substitutions in them. The polypeptide is useful in manufacturing a medicament for treating a syndrome associated with a human disease selected from a pathology associated with the polypeptide. The NOVX polypeptides, polynucleotides and antibodies are useful in treating or preventing NOVX-associated disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's disease, Alzheimer's disease, infections, multiple sclerosis, cancer-associated cachexia, and other wasting disorders associated with chronic diseases. The nucleic acids and polypeptides may also be used as targets for the identification of small molecules that modulate or inhibit e.g. neurogenesis, cell differentiation, cell proliferation, haematopoiesis, wound healing and angiogenesis. In gene therapy, in generation of antibodies that bind immunospecifically to NOVX substances for use in therapeutic or diagnostic methods. The nucleic acids are further used as hybridisation probes, in chromosome mapping, tissue typing, preventive medicine, and pharmacogenomics. The polypeptides are also useful as

CC vaccines. This is the amino acid sequence of a novel human G-protein
CC coupled receptor related protein NOV
XX
XX
SQ Sequence 398 AA;
Query Match 90.0%; Score 618; DB 6; Length 398;
Best Local Similarity 100.0%; Pred. No. 2.8e-62;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDIPINSPELETLTHITTKLAENNAFFYFKIDNVKKARQV 62
DB 224 GKDFVQPTKICVGCPRDIPINSPELETLTHITTKLAENNAFFYFKIDNVKKARQV 283
QY 63 AGKYFIDFVARETTCSKESNEELTESCTKKGSLDCNAEVVVPWEKKIYPTV 118
DB 284 AGKYFIDFVARETTCSKESNEELTESCTKKGSLDCNAEVVVPWEKKIYPTV 339

RESULT 8
ADE76864
ID ADE76864 standard; protein; 427 AA.
XX
XX ADE76864;
XX
XX 29-JAN-2004 (first entry)
XX
XX Human protein expressed in a liver disorder #9.

XX human; liver disorder; hyperlipidaemia; hypertension; type II diabetes;
KW tumour; liver; inflammatory disorder; immune response disorder;
KW high-throughput screening; differential gene expression; gene therapy.
XX
XX Homo sapiens.
XX
XX US2003108871-A1.
XX
XX 12-JUN-2003.
XX
XX 30-JUL-2001; 2001US-00919039.
XX
XX 28-JUL-2000; 2000US-0222113P.
XX
XX (KASE/) KASER M R.
XX
XX Kaser MR;
XX
XX WPI; 2004-031227/03.
XX N-PSDB; ADE76863.

Composition comprising several cDNAs that are differentially expressed in treated human C3A liver cell cultures, useful for treating liver disorders.

Claim 1; SEQ ID NO 29; 41pp; English.

The invention relates to a composition comprising several cDNAs that are differentially expressed in a liver disorder. The composition is useful for treating liver disorder such as hyperlipidaemia, hypertension, type II diabetes, tumours of the liver and disorders of the inflammatory and immune response. The composition is useful for a high-throughput method of screening several molecules or compounds to identify a ligand which specifically binds a cDNA. A protein encoded by the cDNA is useful for a high-throughput method for using a protein to screen several molecules or compounds to identify at least one ligand which specifically binds the protein which involves combining the protein encoded by the cDNA with several of molecules or compounds under conditions to allow specific binding, and detecting specific binding between the protein and a molecule or compound, therefore identifying a ligand which specifically binds the protein. The composition is useful for detecting and quantifying differential gene expression, can be used in gene therapy, to formulate prognosis and to design a treatment regimen and to monitor the efficacy of treatment. The present sequence represents the amino acid sequence of a protein encoded by a cDNA differentially expressed in a

CC liver disorder.
 XX Sequence 427 AA;
 SQ Query Match 90.0%; Score 618; DB 8; Length 427;
 Best Local Similarity 100.0%; Pred. No. 3.1e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPPKICVGCPRDPTNSPELEETLTHITIKLAENNAFFYKIDNVKKARQVW 62
 DB 253 GKDFVQPPKICVGCPRDPTNSPELEETLTHITIKLAENNAFFYKIDNVKKARQVW 312

QY 63 AGKVFIDFVARETTCSKESNEELTESCETKLGSLDCNABVYVVPWEKKIYPTV 118
 DB 313 AGKVFIDFVARETTCSKESNEELTESCETKLGSLDCNABVYVVPWEKKIYPTV 368

RESULT 9
 ABU99144
 ID ABU99144 standard; protein; 615 AA.
 AC ABU99144;
 XX
 DT 01-AUG-2003 (first entry)
 DE Novel human GPCR related protein NOV12b.
 XX Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiac; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX Homo sapiens.
 OS
 XX WO200299116-A2.
 PN
 XX 12-DEC-2002.
 PD
 PF 04-JUN-2002; 2002WO-US017428.
 XX
 PR 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295661P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 14-JUN-2001; 2001US-0296418P.
 PR 15-JUN-2001; 2001US-0298285P.
 PR 21-JUN-2001; 2001US-0298556P.
 PR 26-JUN-2001; 2001US-0299499P.
 PR 28-JUN-2001; 2001US-0300883P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 27-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-0322293P.
 PR 17-SEP-2001; 2001US-0322706P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR 03-JUN-2002; 2002US-00363676.
 XX
 PA (CURA-) CURAGEN CORP.
 XX
 PI Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;
 PI Gangolli EA, Gerlach VU, Gorman L, Guo X, Hjalt T, Kekuda R, Li L;
 PI Macdougall JR, Malyankar UM, Millet I, Padigar M, Patturajan M;
 PI Pena CEA, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CAM;

PI Voss EZ, Zerhusen BD;
 XX WPI; 2003-140627/13.
 DR N-PSDB; ACD03648.
 XX New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 PS Claim 1; Page 144; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-561 residue amino acid sequences, given in the specification, a
 CC mature form of amino acid sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX Sequence 615 AA;
 SQ Query Match 90.0%; Score 618; DB 6; Length 615;
 Best Local Similarity 100.0%; Pred. No. 5.1e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPPKICVGCPRDPTNSPELEETLTHITIKLAENNAFFYKIDNVKKARQVW 62
 DB 224 GKDFVQPPKICVGCPRDPTNSPELEETLTHITIKLAENNAFFYKIDNVKKARQVW 283

QY 63 AGKVFIDFVARETTCSKESNEELTESCETKLGSLDCNABVYVVPWEKKIYPTV 118
 DB 284 AGKVFIDFVARETTCSKESNEELTESCETKLGSLDCNABVYVVPWEKKIYPTV 339

RESULT 10
 ABU78707
 ID ABU78707 standard; protein; 626 AA.
 AC ABU78707;
 XX
 DT 18-JUL-2002 (first entry)
 DE Human high molecular weight kininogen (HK) mature protein SEQ ID NO:1.
 XX Human; kininogen; high molecular weight kininogen; HK; D5 domain;
 KW D5 receptor; angiogenesis; endothelial cell; cytosolic; antitumour;
 KW antiatherosclerotic; vasotropic; vulnery; tranquilliser; thrombolytic;
 KW ophthalmological; gynaecological; antidiabetic; antarthritic;
 KW antiangiogenic; antiapoptotic; endocrine; apoptosis; gene therapy.
 XX Homo sapiens.
 OS
 XX Key Location/Qualifiers
 FT Domain 384..508
 FT /label= D5_domain
 XX WO200214369-A2.
 PN

XX 21-FEB-2002.
 XX 24-JUL-2001; 2001WO-US023185.
 XX 24-JUL-2000; 2000US-0220194P.
 XX (ATTE-) ATTENUON LLC.
 XX Mazar AP, Juarez JC;
 XX WPI; 2002-393611/42.
 XX Novel human kininogen D5 domain polypeptides useful for treating
 XX conditions associated with endothelial cell migration, proliferation,
 XX invasion or angiogenesis, e.g. arthritis, macular degeneration, benign
 XX hyperplasia.
 XX Disclosure; Page 13; 84pp; English.
 XX The present invention describes an isolated polypeptide (I) that
 XX corresponds to the D5 domain of human kininogen, or biologically active
 XX peptide fragment, homologue or functional derivative, and which: (a)
 XX inhibits angiogenesis; (b) binds to the D5 binding site on endothelial
 XX cells (EC); (c) activates signalling pathways leading to the introduction
 XX of apoptosis in EC; and/or (d) inhibits the signalling pathway required
 XX for maintenance of EC viability. (I) has cytostatic, antitumour,
 XX antiatherosclerotic, vasorropic, vulnery, tranquiliser, thrombolytic,
 XX ophthalmological, gynaecological, antiulcer, antidiabetic, antiarthritic,
 XX antiangiogenic, antiapoptotic and endocrine activities. An antibody (IX)
 XX specific for an epitope of (I) is useful for inhibiting tumour growth or
 XX angiogenesis in a subject. (I), a D5 fusion polypeptide (II) or a dimeric
 XX or trimeric fusion polypeptide (III) can be used for inhibiting EC
 XX migration, proliferation, invasion, or angiogenesis, or for inducing EC
 XX apoptosis. An angiogenic EC-targeting pharmaceutical composition (X)
 XX comprising (I), (II), or (III), can be used for treating a subject having
 XX a disease or condition associated with undesired EC migration,
 XX proliferation, invasion or angiogenesis. (I), (II), or (III) can be used
 XX for isolating a D5 domain binding molecule from a complex mixture and for
 XX isolating or enriching cells expressing D5 domain binding sites from a
 XX cell mixture. The present sequence represents the mature human high
 XX molecular weight kininogen (HK) protein, which is given in the
 XX exemplification of the present invention
 XX SQ Sequence 626 AA;
 Query Match 90.0%; Score 618; DB 5; Length 626;
 Best Local Similarity 100.0%; Pred. No. 5.3e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITTKLNAENNAFFYFKIDNVKKARVQV 62
 Db 235 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITTKLNAENNAFFYFKIDNVKKARVQV 294
 QY 63 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 118
 Db 295 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 350
 RESULT 11
 ABG21101
 ID ABG21101 standard; protein; 644 AA.
 XX AC ABG21101;
 XX 18-FEB-2002 (first entry)
 XX Novel human diagnostic protein #21092.
 XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
 XX food supplement; medical imaging; diagnostic; genetic disorder.
 XX Homo sapiens.

XX WO200175067-A2.
 XX 11-OCT-2001.
 XX 30-MAR-2001; 2001WO-US008631.
 XX 31-MAR-2000; 2000US-00540217.
 XX 23-AUG-2000; 2000US-00649167.
 XX (HYSE-) HYSEQ INC.
 XX Drmanac RT, Liu C, Tang YT;
 XX WPI; 2001-639362/73.
 XX N-PSDB; AAS85288.
 XX New isolated polynucleotide and encoded polypeptides, useful in
 XX diagnostics, forensics, gene mapping, identification of mutations
 XX responsible for genetic disorders or other traits and to assess
 XX biodiversity.
 XX Claim 20; SEQ ID NO 51460; 103pp; English.
 XX The invention relates to isolated polynucleotide (I) and polypeptide (II)
 XX sequences. (II) is useful as hybridisation probes, polymerase chain
 XX reaction (PCR) primers, oligomers, and for chromosome and gene mapping.
 XX and in recombinant production of (II). The polynucleotides are also used
 XX in diagnostics as expressed sequence tags for identifying expressed
 XX genes. (I) is useful in gene therapy techniques to restore normal
 XX activity of (II) or to treat disease states involving (II). (II) is
 XX useful for generating antibodies against it, detecting or quantitating a
 XX polypeptide in tissue, as molecular weight markers and as a food
 XX supplement. (II) and its binding partners are useful in medical imaging
 XX of sites expressing (II). (I) and (II) are useful for treating disorders
 XX involving aberrant protein expression or biological activity. The
 XX polypeptide and polynucleotide sequences have applications in
 XX diagnostics, forensics, gene mapping, identification of mutations
 XX responsible for genetic disorders or other traits to assess biodiversity
 XX and to produce other types of data and products dependent on DNA and
 XX amino acid sequences. ABG0010-ABG30377 represent novel human diagnostic
 XX amino acid sequences of the invention. Note: The sequence data for this
 XX patent did not appear in the printed specification, but was obtained in
 XX electronic format directly from WIPO at
 XX ftp.wipo.int/pub/published_pct_sequences
 XX SQ Sequence 644 AA;
 Query Match 90.0%; Score 618; DB 4; Length 644;
 Best Local Similarity 100.0%; Pred. No. 5.5e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITTKLNAENNAFFYFKIDNVKKARVQV 62
 Db 253 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITTKLNAENNAFFYFKIDNVKKARVQV 312
 QY 63 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 118
 Db 313 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 368
 RESULT 12
 ABB78710
 ID ABB78710 standard; protein; 644 AA.
 XX AC ABB78710;
 XX 18-JUL-2002 (first entry)
 XX Human high molecular weight kininogen (HK) protein.
 XX Human; kininogen; high molecular weight kininogen; HK; D5 domain;
 XX D5 receptor; angiogenesis; endothelial cell; cytostatic; antitumour;

KW antiatherosclerotic; vasotropic; vulnerary; tranquilliser; thrombolytic;
 KW ophthalmological; gynaecological; antitumor; antidiabetic; antiarthritic;
 KW antiangiogenic; antiapoptotic; endocrine; apoptosis; gene therapy.

OS Homo sapiens.

XX Key Location/Qualifiers
 FH Peptide 1..18
 FT /label= signal
 FT Protein 19..644
 FT /label= mature_human_high_molecular_weight_kininogen
 FT Disulfide-bond 28..614
 FT Disulfide-bond 83..94
 FT Disulfide-bond 107..126
 FT Disulfide-bond 142..145
 FT Disulfide-bond 206..218
 FT Disulfide-bond 229..248
 FT Disulfide-bond 264..267
 FT Disulfide-bond 328..340
 FT Disulfide-bond 351..370
 FT Domain 402..526
 FT /label= D5_domain

WO200214369-A2.

21-FEB-2002.

24-JUL-2001; 2001WO-US023185.

24-JUL-2000; 2000US-0220194P.

(ATTE-) ATTENDON LLC.

Mazar AP, Juarez JC;

WPI; 2002-393611/42.

Novel human kininogen D5 domain polypeptides useful for treating conditions associated with endothelial cell migration, proliferation, invasion or angiogenesis, e.g. arthritis, macular degeneration, benign hyperplasia.

Disclosure; Fig 1B-E; 84pp; English.

The present invention describes an isolated polypeptide (I) that corresponds to the D5 domain of human kininogen, or biologically active peptide fragment, homologue or functional derivative, and which: (a) inhibits angiogenesis; (b) binds to the D5 binding site on endothelial cells (EC); (c) activates signalling pathways leading to the introduction of apoptosis in EC; and/or (d) inhibits the signalling pathway required for maintenance of EC viability. (I) has cytostatic, antitumor, antiatherosclerotic, vasotropic, vulnerary, tranquilliser, thrombolytic, ophthalmological, gynaecological, antitumor, antidiabetic, antiarthritic, antiangiogenic, antiapoptotic and endocrine activities. An antibody (IX) specific for an epitope of (I) is useful for inhibiting tumour growth or angiogenesis in a subject. (I), a D5 fusion polypeptide (II) or a dimeric or trimeric fusion polypeptide (III) can be used for inhibiting EC migration, proliferation, invasion, or angiogenesis, or for inducing EC apoptosis. An angiogenic EC-targeting pharmaceutical composition (X) comprising (I), (II), or (III), can be used for treating a subject having a disease or condition associated with undesired EC migration, proliferation, invasion or angiogenesis. (I), (II), or (III) can be used for isolating a D5 domain binding molecule from a complex mixture and for isolating or enriching cells expressing D5 domain binding sites from a cell mixture. The present sequence represents the human high molecular weight kininogen (HK) protein, which is given in the exemplification of the present invention

Sequence 644 AA;

Query Match 90.0%; Score 618; DB 5; Length 644;

Best Local Similarity 100.0%; Pred. No. 5.5e-62;

Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDPTNSPELEETLTHITIKLNAENATFYFKIDNVKKARQVQV 62
 DB 253 GKDFVQPTKICVGCPRDPTNSPELEETLTHITIKLNAENATFYFKIDNVKKARQVQV 312
 QY 63 AGKKYFIDFVARETTCSKESNEELTESCTKKLGSLDCNAEYVVVPWEKKIYPTV 118
 DB 313 AGKKYFIDFVARETTCSKESNEELTESCTKKLGSLDCNAEYVVVPWEKKIYPTV 368

RESULT 13

ABU99150

ID ABU99150 standard; protein; 644 AA.

AC ABU99150;

XX 01-AUG-2003 (first entry)

Novel human GPCR related protein NOV12h.

KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytostatic; cardiact; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.

XX Homo sapiens.

XX WO200299116-A2.

XX 12-DEC-2002.

XX 04-JUN-2002; 2002WO-US017428.

XX 04-JUN-2001; 2001US-0295607P.

XX 04-JUN-2001; 2001US-0295661P.

XX 06-JUN-2001; 2001US-0298404P.

XX 06-JUN-2001; 2001US-0298418P.

XX 14-JUN-2001; 2001US-0298285P.

XX 15-JUN-2001; 2001US-0298556P.

XX 21-JUN-2001; 2001US-0299949P.

XX 26-JUN-2001; 2001US-0300883P.

XX 28-JUN-2001; 2001US-0301550P.

XX 13-AUG-2001; 2001US-0311972P.

XX 27-AUG-2001; 2001US-0315071P.

XX 29-AUG-2001; 2001US-0315660P.

XX 14-SEP-2001; 2001US-0322293P.

XX 17-SEP-2001; 2001US-0322706P.

XX 14-DEC-2001; 2001US-0341186P.

XX 28-FEB-2002; 2002US-0361189P.

XX 12-MAR-2002; 2002US-0363673P.

XX 12-MAR-2002; 2002US-0363676P.

XX 03-JUN-2002; 2002US-00363676.

(CURA-) CURAGEN CORP.

XX Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;

XX Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalt T, Kekuda R, Li L;

XX Macdougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M; CAW;

XX Pena CE, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet

XX Voss EZ, Zerhusen BD;

XX WPI; 2003-140627/13.

XX DR N-PSDB; ACD03654.

XX New NOVX polypeptides and nucleic acids, useful for preventing or

XX treating NOVX-associated disorders, e.g. cancer, cardiomyopathy.


```

Db      313 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVYVWPWEKKIYPTV 368
RESULT 15
AAB37447
ID      AAB37447 standard; protein; 122 AA.
XX
XX      AC      AAB37447;
XX      DT      21-FEB-2001 (first entry)
XX      DE      Human kininogen D3.
XX      KW      Enzyme; legumain; endopeptidase; cystatin; human; kininogen.
XX      OS      Homo sapiens.
XX      PN      WO2000064945-A1.
XX      PD      02-NOV-2000.
XX      PF      20-APR-2000; 2000WO-GB001571.
XX      PR      22-APR-1999; 99GB-00009133.
XX      PA      (BABR-) BABRAHAM INST.
XX      PI      Abrahamson M, Barrett AJ;
XX      DR      WPI; 2000-687316/67.
XX      PT      Inhibition of mammalian legumain or legumain-related endopeptidase by
AP      cystatin involves interaction with second papain-non-reactive site of
XX      PS      cystatin.
XX      SQ      Disclosure; Fig 4; 45pp; English.
XX
XX      The present invention relates to inhibition of the enzymatic activity of
CC      legumain or a legumain-related endopeptidase by cystatin. The inhibition
CC      involves an interaction between legumain and a papain-non-reactive site
CC      of cystatin. Legumain (EC 3.4.22.34) is a cysteine endopeptidase, and
CC      performs a protein-processing function. The present sequence is human
CC      kininogen D3, which was used in the present invention. Kininogen is a
CC      type 3 cystatin
XX
XX      Query Match      85.3%; Score 586; DB 3; Length 122;
XX      Best Local Similarity 100.0%; Pred. No. 2.7e-59;
XX      Matches 110; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY      9 PPTKICVGCPRDIPNPSPELETLTHTITKLNANNATFYFKIDNVKKARVQVVGKKYF 68
Db      1 PPTKICVGCPRDIPNPSPELETLTHTITKLNANNATFYFKIDNVKKARVQVVGKKYF 60
QY      69 IDFVARETTCSKESNEELTESCETKLGSLDCNAEVVYVWPWEKKIYPTV 118
Db      61 IDFVARETTCSKESNEELTESCETKLGSLDCNAEVVYVWPWEKKIYPTV 110

Search completed: September 24, 2004, 14:08:38
Job time : 52.308 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:07:01 ; Search time 14,732 Seconds
(without alignments)
445.051 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687

Sequence: 1 GSGKDFVQPTKICVGPDR.....VPWEKKIYPTVTWNHECEF 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents, AA.*
1: /cgn2_6/ptodata/2/iaa/5A_COMB.pep.*
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3: /cgn2_6/ptodata/2/iaa/6A_COMB.pep.*
4: /cgn2_6/ptodata/2/iaa/6B_COMB.pep.*
5: /cgn2_6/ptodata/2/iaa/PCTUS_COMB.pep.*
6: /cgn2_6/ptodata/2/iaa/backfiles.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	558	81.2	117	1	US-08-193-114B-1
2	556.5	81.0	117	5	PCT-US92-06809-1
3	163.5	23.8	145	2	US-08-832-535-2
4	163.5	23.8	145	3	US-09-019-485-2
5	163.5	23.8	145	3	US-09-019-485-3
6	163.5	23.8	145	3	US-09-431-480-9
7	163.5	23.8	145	3	US-09-617-302-9
8	163.5	23.8	145	4	US-09-528-436B-2
9	163	23.7	178	2	US-08-791-522-1
10	163	23.7	178	3	US-08-314-777-1
11	138.5	20.2	121	4	US-09-775-932-14
12	138.5	20.2	128	4	US-09-775-932-12
13	138.5	20.2	149	2	US-08-461-030C-2
14	138.5	20.2	149	3	US-08-744-138-2
15	138.5	20.2	149	3	US-09-431-480-8
16	138.5	20.2	149	3	US-09-431-480-10
17	138.5	20.2	149	3	US-09-617-302-8
18	138.5	20.2	149	3	US-09-617-302-10
19	138.5	20.2	149	4	US-09-241-376-2
20	138.5	20.2	149	4	US-09-940-497-2
21	137.5	19.9	112	4	US-08-849-303-16
22	136.5	19.9	118	4	US-09-775-932-24
23	135.5	19.7	146	6	5432264-6
24	134	18.5	148	5	PCT-US95-07135-2
25	132.5	19.3	120	4	US-09-775-932-2
26	132.5	19.3	145	2	US-08-832-535-11
27	132.5	19.3	146	2	US-08-791-522-3

28 132.5 19.3 146 3 US-08-744-138-3
29 132.5 19.3 146 3 US-09-019-485-4
30 132.5 19.3 146 3 US-09-314-777-3
31 132.5 19.3 146 3 US-09-431-480-6
32 132.5 19.3 146 3 US-09-617-302-6
33 132.5 19.3 146 4 US-09-241-376-3
34 132.5 19.3 146 4 US-09-528-436B-3
35 132.5 19.3 146 4 US-09-886-319A-47
36 132.5 19.3 146 4 US-09-940-497-3
37 132.5 19.3 146 4 US-09-976-594-37
38 132.5 19.3 146 4 US-08-849-303-17
39 132.5 19.3 146 5 PCT-US95-07135-9
40 132 19.3 26 3 US-08-676-242-15
41 131.5 19.1 382 4 US-09-599-360B-93
42 130 18.9 127 4 US-08-849-303-19
43 129.5 18.9 140 4 US-09-886-319A-46
44 129.5 18.9 140 4 US-09-886-319A-48
45 128 18.6 111 4 US-08-849-303-26

ALIGNMENTS

RESULT 1
US-08-193-114B-1
; Sequence 1, Application US/08193114B
; Patent No. 5472945
; GENERAL INFORMATION:
; APPLICANT: Schmaier, Alvin H.
; APPLICANT: Jiang, Yongping
; TITLE OF INVENTION: Modulation of Blood
; TITLE OF INVENTION: Pressure and Inhibition of Platelet Activation
; TITLE OF INVENTION: with Kininogen Fragment
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Seidel, Gonda, Lavorgna &
; ADDRESSEE: Monaco, P.C.
; STREET: 1800 Two Penn Center Plaza
; CITY: Philadelphia
; STATE: Pennsylvania
; COUNTRY: U.S.A.
; ZIP: 19102
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb
; COMPUTER: IBM PS/2
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: WordPerfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/193,114B
; FILING DATE: 9 February 1994
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: U.S. Application
; APPLICATION NUMBER: Serial No. 5472945 07/744,545
; FILING DATE: 13 August 1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Monaco, Daniel A.
; REGISTRATION NUMBER: 30,480
; REFERENCE/DOCKET NUMBER: 6056-137 CII
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-8383
; TELEFAX: (215) 568-5549
; TELEX: No. 5472945e
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 117 amino acids
; TYPE: peptide
; TOPOLOGY: linear
US-08-193-114B-1

Query Match 81.2%; Score 558; DB 1; Length 117;
Best Local Similarity 100.0%; Pred. No. 2,3e+55;
Matches 105; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 14 CVGCPDRIPNSPELEETLTHITKLNAAENATFYFKIDNVKKARVQVWAGKYYFIDFVA 73
Db 1 CVGCPDRIPNSPELEETLTHITKLNAAENATFYFKIDNVKKARVQVWAGKYYFIDFVA 60

QY 74 RETTCSKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTV 118
Db 61 RETTCSKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTV 105

RESULT 2
PCT-US92-06809-1
; Sequence 1, Application PC/TUS9206809
; GENERAL INFORMATION:
; APPLICANT: Schmaier, Alvin H.
; TITLE OF INVENTION: Modulation of Blood
; TITLE OF INVENTION: Pressure by Altering Bradykinin Levels
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Temple University - Of the
; ADDRESS: Commonwealth System of Higher Education
; STREET: 406 University Services
; STREET: Building
; CITY: Philadelphia
; STATE: Pennsylvania
; COUNTRY: U.S.A.
; ZIP: 19122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb
; COMPUTER: IBM PS/2
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: WordPerfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US92/06809
; FILING DATE: 19910813
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: U.S. Application
; APPLICATION NUMBER: Serial No. 744,545
; FILING DATE: 13 August 1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Monaco, Daniel A.
; REGISTRATION NUMBER: 30,480
; REFERENCE/DOCKET NUMBER: 6056-137
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-8383
; TELEFAX: (215) 568-5549
; TELEX:
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 117 amino acids
; TYPE: AMINO ACID
; TOPOLOGY: linear
PCT-US92-06809-1
Query Match 81.0%; Score 556.5; DB 5; Length 117;
Best Local Similarity 93.8%; Pred. No. 3.3e-55;
Matches 106; Conservative 1; Mismatches 1; Indels 5; Gaps 1;

QY 14 CVGCPDRIPNSPELEETLTHITKLNAAENATFYFKIDNVKKARVQVWAGKYYFIDFVA 73
Db 1 CVGCPDRIPNSPELEETLTHITKLNAAENATFYFKIDNVKKARVQVWAGKYYFIDFVA 60

QY 74 RETTCSKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTV 126
Db 61 RETTCSKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTV 108

RESULT 3
US-08-832-535-2
; Sequence 2, Application US/08832535
; Patent No. 5919658
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; GENERAL INFORMATION:
; APPLICANT: NI, JIAN
; APPLICANT: LI, HAODONG
; APPLICANT: YU, GUO-LIANG
; APPLICANT: GENTZ, REINER L
; TITLE OF INVENTION: HUMAN CYSTATIN F
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HUMAN GENOME SCIENCES, INC.
; STREET: 9410 KEY WEST AVENUE
; CITY: ROCKVILLE
; STATE: MD
; COUNTRY: US
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/832,535
; FILING DATE: 03-APR-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: KIMBALL, PAUL C.
; REGISTRATION NUMBER: 34,610
; REFERENCE/DOCKET NUMBER: PF265
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (201) 994-1700
; TELEFAX: (201) 994-1744
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-832-535-2
Query Match 23.8%; Score 163.5; DB 2; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVCPDRIPNSPELEETLTHITKLNAAENATFYFKIDNVKKARVQVWAGKYYFID 70
Db 32 SRVPGFPKTIKTNDGVLQAAARYSVEKNNTNDMFLFKESRITRALVQIVKGLKYMLE 91

QY 71 FVARETTCSKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTV 124
Db 92 VEIGRTTCKKNQHLRL-DDCDFQTNHTLKQTLSCYSEVWVWVW-----LQHF 138

RESULT 4
US-09-019-485-2
; Sequence 2, Application US/09019485
; Patent No. 6066617
; GENERAL INFORMATION:
; APPLICANT: Li, Haodong
; APPLICANT: Yu, Guo-liang
; APPLICANT: Gentz, Reiner
; APPLICANT: Ni, Jian
; TITLE OF INVENTION: Cystatin F
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: MD
; COUNTRY: US
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
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; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/019,485
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Benson, Robert H.
; REGISTRATION NUMBER: 30,446
; REFERENCE/DOCKET NUMBER: PF265P1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 3013098504
; TELEFAX: 3013098439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-019-485-2

Query Match 23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 3; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVGCPRDIPNTSPLEETLTHTITKLAENNATYFKIDNVKKARQVQVAGKKYFID 70
   :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: ::
Db 32 SRVKGPGFKTIKNDPGVQAARYSVEKFNNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
   :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: ::

QY 71 FVARETTCSKESNEELTESCE--TKLQSGLDCAEVVYVYVPWKKIYPTVTVAHWE 124
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 92 VEIGRTTKCKNQHQURL-DDCDFQTNHTLKQTLSCYSEVVVVPW-----LQHFE 138
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 5
US-09-019-485-3
; Sequence 3, Application US/09019485
; Patent No. 6066617
; GENERAL INFORMATION:
; APPLICANT: Li, Haodong
; APPLICANT: Yu, Guo-Liang
; APPLICANT: Gentz, Reiner
; APPLICANT: Ni, Jian
; TITLE OF INVENTION: Cystatin F
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: MD
; COUNTRY: US
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/019,485
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Benson, Robert H.
; REGISTRATION NUMBER: 30,446
; REFERENCE/DOCKET NUMBER: PF265P1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 3013098504
; TELEFAX: 3013098439
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear

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; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 12
; LENGTH: 128
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-12

Query Match      20.2%; Score 138.5; DB 4; Length 128;
Best Local Similarity 31.5%; Pred. No. 5.2e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPITNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWVAGKYYFI 69
Db 9 POERWVGBELRLSDPDPQVQKAAQAAVASVYNGNSIYFRDTHIIKAQSQLVAGIKYFL 68
QY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
Db 69 TWEMGSTDCRTRVTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 115

RESULT 13
US-08-461-030C-2
; Sequence 2, Application US/08461030C
; Patent No. 5985601
; GENERAL INFORMATION:
; APPLICANT: Ni, Jian
; APPLICANT: Yu, Guo-Liang
; APPLICANT: Gentz, Reiner
; APPLICANT: Rosen, Craig A.
; TITLE OF INVENTION: Human Cystatin B
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Ave
; CITY: Rockville
; STATE: MD
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/461,030C
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: A. Anders, Brookes
; REGISTRATION NUMBER: 36,373
; REFERENCE/DOCKET NUMBER: PF202
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 301-301-8504
; TELEFAX: 301-309-8439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 149 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-461-030C-2

Query Match      20.2%; Score 138.5; DB 2; Length 149;
Best Local Similarity 31.5%; Pred. No. 6.4e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPITNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWVAGKYYFI 69
Db 30 POERWVGBELRLSDPDPQVQKAAQAAVASVYNGNSIYFRDTHIIKAQSQLVAGIKYFL 89

; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 178 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: 30443
US-09-314-777-1

Query Match      23.7%; Score 163; DB 3; Length 178;
Best Local Similarity 34.0%; Pred. No. 1.4e-10;
Matches 35; Conservative 20; Mismatches 44; Indels 4; Gaps 2;

QY 11 TKICVGCPRDIPITNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWVAGKYYFI 70
Db 54 SRVKGFPKTIKNDPGVLQAARYSVERFNCTNDMLFKESRITRALVQIVKGLKYLE 113
QY 71 FVARETTCSKESNEELTSCSE---TKLQGLSDCNAEVVVPW 110
Db 114 VEIGRTCKKNQHLRL-DCDFQTNHTLKQTLSCYSEVVVVPW 155

RESULT 11
US-09-775-932-14;
; Sequence 14, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 14
; LENGTH: 121
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-14

Query Match      20.2%; Score 138.5; DB 4; Length 121;
Best Local Similarity 31.5%; Pred. No. 4.9e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPITNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWVAGKYYFI 69
Db 2 POERWVGBELRLSDPDPQVQKAAQAAVASVYNGNSIYFRDTHIIKAQSQLVAGIKYFL 61
QY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
Db 62 TWEMGSTDCRTRVTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 108

RESULT 12
US-09-775-932-12
; Sequence 12, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05

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; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 12
; LENGTH: 128
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-12

Query Match      20.2%; Score 138.5; DB 4; Length 128;
Best Local Similarity 31.5%; Pred. No. 5.2e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPITNSPELEETLTHITIKLNAENNATFYFKIDNVKKARVQVWVAGKYYFI 69
Db 9 PQERWVGBELRLSDPDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSQLVAGIKYFL 68
QY 70 DFVARETTCSKE-----SNEELTESCETKKLGQ--SLDCNAEYVVPWE 111
Db 69 TWEMGSTDCRTRVTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 115

RESULT 13
US-08-461-030C-2
; Sequence 2, Application US/08461030C
; Patent No. 5985601
; GENERAL INFORMATION:
; APPLICANT: Ni, Jian
; APPLICANT: Yu, Guo-Liang
; APPLICANT: Gentz, Reiner
; APPLICANT: Rosen, Craig A.
; TITLE OF INVENTION: Human Cystatin B
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Ave
; CITY: Rockville
; STATE: MD
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/461,030C
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: A. Anders, Brookes
; REGISTRATION NUMBER: 36,373
; REFERENCE/DOCKET NUMBER: PF202
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 301-301-8504
; TELEFAX: 301-309-8439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 149 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-461-030C-2

Query Match      20.2%; Score 138.5; DB 2; Length 149;
Best Local Similarity 31.5%; Pred. No. 6.4e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPITNSPELEETLTHITIKLNAENNATFYFKIDNVKKARVQVWVAGKYYFI 69
Db 30 PQERWVGBELRLSDPDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSQLVAGIKYFL 89

; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 178 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: 30443
US-09-314-777-1

Query Match      23.7%; Score 163; DB 3; Length 178;
Best Local Similarity 34.0%; Pred. No. 1.4e-10;
Matches 35; Conservative 20; Mismatches 44; Indels 4; Gaps 2;

QY 11 TKICVGCPRDIPITNSPELEETLTHITIKLNAENNATFYFKIDNVKKARVQVWVAGKYYFI 70
Db 54 SRVKGFPKTIKNDPGVLQAARYSVERFNCTNDMLFKESRITRALVQIVKGLKYLE 113
QY 71 FVARETTCSKESNEELTSCSE---TKKLGQSLDCNAEYVVPW 110
Db 114 VEIGRTCKKNQHLRL-DCDFQTNHTLKQTLSCYSEVVVVPW 155

RESULT 11
US-09-775-932-14;
; Sequence 14, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 14
; LENGTH: 121
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-14

Query Match      20.2%; Score 138.5; DB 4; Length 121;
Best Local Similarity 31.5%; Pred. No. 4.9e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPITNSPELEETLTHITIKLNAENNATFYFKIDNVKKARVQVWVAGKYYFI 69
Db 2 PQERWVGBELRLSDPDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSQLVAGIKYFL 61
QY 70 DFVARETTCSKE-----SNEELTESCETKKLGQ--SLDCNAEYVVPWE 111
Db 62 TWEMGSTDCRTRVTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 108

RESULT 12
US-09-775-932-12
; Sequence 12, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05

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QY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
Db 90 TWMGSTDCKRTRVTGDHVDLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 136

RESULT 14

US-08-744-138-2

; Sequence 2, Application US/08744138

; Patent No. 6011012

; GENERAL INFORMATION:

; APPLICANT: Gentz, Reiner L.

; APPLICANT: Ni, Jian

; APPLICANT: Rosen, Craig A.

; APPLICANT: Yu, Guo-Liang

; TITLE OF INVENTION: Human Cystatin E

; NUMBER OF SEQUENCES: 13

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Human Genome Sciences, Inc.

; STREET: 9410 Key West Avenue

; CITY: Rockville

; STATE: Maryland

; COUNTRY: USA

; ZIP: 20850

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent In Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/744,138

; FILING DATE:

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: Brookes, A. Anders

; REGISTRATION NUMBER: 36,373

; REFERENCE/DOCKET NUMBER: PF202P1

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 301 309 8504

; TELEFAX: 301 309 8512

; INFORMATION FOR SEQ ID NO: 2:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 149 amino acids

; TYPE: amino acid

; TOPOLOGY: linear

; MOLECULE TYPE: protein

; US-08-744-138-2

Query Match 20.2%; Score 138.5; DB 3; Length 149;
Best Local Similarity 31.5%; Pred. No. 6.4e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPITNSPELEETLTHITIKLNAENNATFYFKIDNVKKARVQVWAGKKYFI 69

Db 30 PQERMVGEIRDLSPPDPQVQKAAQAAVSYNMGNSIYFRDTHIIKAQSQLVAGIKYFL 89

QY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111

Db 90 TWMGSTDCKRTRVTGDHVDLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 136

RESULT 15

US-09-431-480-8

; Sequence 8, Application US/09431480

; Patent No. 6235708

; GENERAL INFORMATION:

; APPLICANT: Holloway, James L.

; APPLICANT: Feldhaus, Andrew

; TITLE OF INVENTION: TESTIS SPECIFIC CYSTATIN-LIKE PROTEIN CYSTATIN T

; FILE REFERENCE: 98-72

; CURRENT APPLICATION NUMBER: US/09/431,480

; CURRENT FILING DATE: 1999-11-01

; EARLIER APPLICATION NUMBER: 60/109,217

; EARLIER FILING DATE: 1998-11-20

; EARLIER APPLICATION NUMBER: 60/156,382
; EARLIER FILING DATE: 1999-09-28
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 149
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-431-480-8

Query Match 20.2%; Score 138.5; DB 3; Length 149;
Best Local Similarity 31.5%; Pred. No. 6.4e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPITNSPELEETLTHITIKLNAENNATFYFKIDNVKKARVQVWAGKKYFI 69

Db 30 PQERMVGEIRDLSPPDPQVQKAAQAAVSYNMGNSIYFRDTHIIKAQSQLVAGIKYFL 89

QY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111

Db 90 TWMGSTDCKRTRVTGDHVDLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 136

Search completed: September 24, 2004, 14:11:37

Job time : 15.732 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:05:18 ; Search time 36.576 Seconds
(without alignments)
1095.549 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687
Sequence: 1 GSKGDFVQPTKICVGCPRD.....VPWEKKIYPTVTVNHWECEF 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL 25:*

1: sp_arChaea:*

2: sp_bacteria:*

3: sp_fungi:*

4: sp_human:*

5: sp_invertebrate:*

6: sp_mammal:*

7: sp_mhc:*

8: sp_organelle:*

9: sp_phages:*

10: sp_plant:*

11: sp_rodent:*

12: sp_virus:*

13: sp_vertebrate:*

14: sp_unclassified:*

15: sp_virus:*

16: sp_bacteriap:*

17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	383	55.7	140	6 Q7YRP6	Q7YRP6 sus scrofa
2	381	55.5	423	11 P70517	P70517 rattus norv
3	378	55.0	430	11 Q63581	Q63581 rattus norv
4	171.5	25.0	167	11 Q5QWL5	Q5QWL5 mus musculu
5	163.5	23.8	167	4 Q7Z4J8	Q7Z4J8 homo sapien
6	152.5	22.2	462	13 Q7ZY91	Q7ZY91 xenopus lae
7	152.5	22.2	462	13 Q7SVH2	Q7SVH2 xenopus lae
8	152.5	22.2	465	13 Q801E5	Q801E5 xenopus lae
9	131.5	19.1	140	11 Q9EPX9	Q9EPX9 mus musculu
10	123.5	18.0	455	11 Q800S8	Q800S8 brachydanio
11	119	17.3	388	11 Q8CB17	Q8CB17 mus musculu
12	117.5	17.1	464	13 Q801Z5	Q801Z5 cyprinus ca
13	113.5	16.5	148	5 Q9NH95	Q9NH95 litomosoid
14	113	16.4	140	11 Q80Y72	Q80Y72 mus musculu
15	111	16.2	146	11 Q8K397	Q8K397 mus musculu
16	111	16.2	149	11 Q9D1B1	Q9D1B1 mus musculu

17	108.5	15.8	112	13 Q98SR4	Q98SR4 acipenser s
18	108.5	15.8	112	13 Q98SR3	Q98SR3 acipenser s
19	107	15.6	139	5 Q9TY2	Q9TY2 caenorhabdi
20	106	15.4	300	13 Q801Z6	Q801Z6 cyprinus ca
21	105	15.3	109	5 Q9TY65	Q9TY65 onchocerca
22	105	15.3	127	5 P90698	P90698 brugia mala
23	104.5	15.2	149	11 Q8VHC1	Q8VHC1 rattus norv
24	104.5	15.2	161	5 Q16159	Q16159 brugia mala
25	102.5	14.9	127	5 Q9U9A1	Q9U9A1 onchocerca
26	101	14.7	148	11 Q9JMH4	Q9JMH4 mus musculu
27	99	14.4	110	10 Q8SA65	Q8SA65 sandersonia
28	98.5	14.3	107	5 Q8T0Y2	Q8T0Y2 sarcophaga
29	98.5	14.3	125	5 Q25620	Q25620 onchocerca
30	97.5	14.2	134	10 Q41825	Q41825 zea mays (m
31	95.5	13.9	143	5 Q61973	Q61973 caenorhabdi
32	94	13.7	122	5 Q44396	Q44396 haemonchus
33	93	13.5	138	4 Q8WXU6	Q8WXU6 homo sapien
34	92	13.4	157	5 Q17108	Q17108 acanthochei
35	90	13.1	139	11 Q8K5A3	Q8K5A3 rattus norv
36	88.5	12.9	92	10 Q9FXN6	Q9FXN6 arabidopsis
37	88.5	12.9	116	10 Q8XS7	Q8XS7 arabidopsis
38	88.5	12.9	124	10 Q41906	Q41906 arabidopsis
39	88.5	12.9	125	10 Q22202	Q22202 arabidopsis
40	88.5	12.9	134	10 P93627	P93627 zea mays (m
41	88.5	12.9	134	10 Q41897	Q41897 zea mays (m
42	88	12.8	199	10 Q9Z270	Q9Z270 brassica ca
43	87.5	12.7	141	11 Q9DAP1	Q9DAP1 mus musculu
44	86.5	12.6	141	11 Q80ZNS	Q80ZNS mus musculu
45	86.5	12.6	349	6 Q14502	Q14502 cercopithec

ALIGNMENTS

RESULT 1

Q7YRP6 ID Q7YRP6 PRELIMINARY; PRT; 140 AA.

AC Q7YRP6; 01-OCT-2003 (TrEMBLrel. 25, Created)

DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)

DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)

DE Low molecular weight kininogen (Fragment).

GN KNG.

OS Sus scrofa (Pig).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.

OX NCBI_TaxID=9823;

RN [1]

RP SEQUENCE FROM N.A.

RA Vonnahme K.A., Fernando S.C., Ross J.A., Ashworth M.D., DeSilva U.,

RA Malayer J.R., Geisert R.D.;

RT "Porcine Endometrial and Conceptus Expression of Kininogens and Plasma

RT Kallikrein in Cyclic and Pregnant Gilts."

RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.

DR EMBL; AY321363; AAP85260.1; -

FT NON_TER 1

FT NON_TER 140

SQ SEQUENCE 140 AA; 15650 MW; 177837836603F777 CRC64;

Query Match 55.7%; Score 383; DB 6; Length 140;

Best Local Similarity 78.9%; Pred. No. 2.8e-30;

Matches 75; Conservative 5; Mismatches 15; Indels 0; Gaps 0;

Qy 24 NSPELEETLTHTTTKLNAENNAATFYFKINDVKARVQVWAGKYFDVFARETTCSEKN 83

Db 1 DSPDLEPLNHSIAKLNAENNAVFYFKIGVEKATVQVWAGKYIVFTARETTCQSN 60

Qy 84 EELTESCETKLGQSLDCNAEVVVPWEKKIYPTV 118

Db 61 EELTESCEIKKPGQILKCNASVVPWEKKIYPTV 95

RESULT 2

Anderson K.P., Croyle M.L., Lingrel J.B.;
"Primary structure of a gene encoding rat T-kininogen.";
Gene 81:115-128(1989).
DR ENBL; M29090; AAA42251.1; JOINED.
DR ENBL; M29083; AAA42251.1; JOINED.
DR ENBL; M29084; AAA42251.1; JOINED.
DR ENBL; M29091; AAA42251.1; JOINED.
DR ENBL; M29085; AAA42251.1; JOINED.
DR ENBL; M29086; AAA42251.1; JOINED.
DR ENBL; M29087; AAA42251.1; JOINED.
DR ENBL; M29088; AAA42251.1; JOINED.
DR ENBL; M29089; AAA42251.1; JOINED.
DR PIR; S68034; S68034.
DR PIR; S68035; S68035.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 3.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
DR SEQUENCE 430 AA; 47618 MW; 45508DEF4BDC978C CRC64;

Query Match 55.0%; Score 378; DB 11; Length 430;
Best Local Similarity 62.1%; Pred.No.3.le-29;
Matches 72; Conservative 13; Mismatches 31; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVCPDRIPNSPELEETLTHITKLNENNAATPVFKIDNVKARVQV 62
Db 252 GUDLFLPKRCRCPRPEIPVDSPELKEALGHSIAQNAQHNIFFFKIDTVKKATSQW 311

QY 63 AGKYFIDFVARETTCSEKNEELTESCETKLGQSLDCAEYVYVPVEKKIYPTV 118
Db 312 AGVIIVIEFIARETNCQSKQSTELTADCEPKHGLQSLNCANVYMPENKVVPTV 367

RESULT 4
Q9QWL5 PRELIMINARY; PRT; 167 AA.
ID Q9QWL5 AC Q9QWL5
DT 01-WAY-2000 (TReMBLrel. 13, Created)
DT 01-WAY-2000 (TReMBLrel. 13, Last sequence update)
DT 01-JUN-2003 (TReMBLrel. 24, Last annotation update)
DE Murine CMAP (CYSTATIN F) (LEUKOCYSTATIN).
GN MURINE CMAP OR CST7.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA Morita M., Arakawa H., Yoshiuchi N.;
RT "A novel cystatin-like metastasis associated gene";
RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
RP [2]
RP SEQUENCE FROM N.A.
RC STRAIN=CSFBL/6J; TISSUE=Embryo;
RX MEDLINE=21083660; PubMed=11721851;
RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RA Aizawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
RA Arakawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
RA Saïto T., Okazaki Y., Gotohori T., Bono H., Kasukawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
RA Fletschmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
RA Kueli P., Lewis S., Matsuo Y., Nikaïdo I., Pesole G., Quackenbush J.,
RA Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawai H., Kohtsuki S.,

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RA Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690(2001).
DR EMBL; AB015224; BAA34940.1; -.
DR EMBL; AK004420; BAB23298.1; -.
DR HSSP; P01034; 1G96.
DR MGD; MGI:1298217; Cst7.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 2.
DR SEQUENCE 167 AA; 18847 MW; 61F776D8445095FE CRC64;

Query Match 25.0%; Score 171.5; DB 11; Length 167;
Best Local Similarity 35.5%; Pred. No. 3.4e-09;
Matches 39; Conservative 22; Mismatches 42; Indels 7; Gaps 3;

QY 4 KDFVQPTKICVGCPRIPITNSPELEETLTHITIKLAENNAFYFKIDNVKKARQVQVVA 63
DB 50 KDLI---SSVKPGFPKTIETNPGVLKAARHSVEKFNCTNDIFLFKESHVSKALVQVVK 106
QY 64 GKYPIDFVARETTCKESNEELTESCE---TKLGGSLDCNAEYVVPW 110
DB 107 GLKTMLEVKIGRTTCRKTMMHQL-DNCDFOINPALKXETLYCYSEVWVWP 155

RESULT 5
Q724J8 PRELIMINARY; PRT; 167 AA.
AC Q724J8
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin F (Leukocystatin).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Carnivora; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Kainine N., Chen X., Rolfs A., Halleck A., Hines L., Eisenstein S.,
RA Koundinya M., Raphael J., Moreira D., Kelley T., LaBaer J., Lin Y.,
RA Phelan M., Farmer A.;
RL Submitted (AUG-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BT009825; AAP88827.1; -.
SQ SEQUENCE 167 AA; 18857 MW; E339025ASBD50177 CRC64;

Query Match 23.8%; Score 163.5; DB 4; Length 167;
Best Local Similarity 31.6%; Pred. No. 2.1e-08;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVGCPRDPTNSPELEETLTHITIKLAENNAFYFKIDNVKKARQVQVQVAGKYFID 70
DB 54 SRVKFGFPKTIKTNDPGVLOARYSVEKFNCTNDIMFLFKESRITRALVQIVKGLKYLE 113
QY 71 FVARETTCKESNEELTESCE---TKLGGSLDCNAEYVVPWPKIYPTVVAHWE 124
DB 114 VEIGRTCKKQCHLRL-DDCFQTHILKQILSCYSEVWVWP-----LQHPF 160

RESULT 6
Q7ZY91 PRELIMINARY; PRT; 462 AA.
AC Q7ZY91
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Similar to fetuin B.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;

RA SEQUENCE FROM N.A.
TISSUE=Embryo;
RA Klein S., Strausberg R.;
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC043891; AAH43891.1; -.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; CY; 2.
DR SEQUENCE 462 AA; 53185 MW; D7BAD339961739FB CRC64;

Query Match 22.2%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 8.4e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

QY 10 PTKICVGCPRDPTNSPELEETLTHITIKLAENNAFYFKIDNVKKARQVQVQVAGK 65
DB 142 PGVILSTCP-DCTANEETPTITETADTLIAEYKNSNTTRYFKIDHIERVRSQWVGP 200
QY 66 KYFIDFVARETTCKESNEELTESC 90
DB 201 SYPIQFTIKETDCMKTQENWLSNC 225

RESULT 7
Q7SYH2 PRELIMINARY; PRT; 462 AA.
AC Q7SYH2
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin domain fetuin-like protein.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
TISSUE=Ventral midgut;
RA Costa R.M.B., Mason J., Lee M., Amaya E., Zorn A.M.;
RA "Novel gene expression domains reveal early patterning of the Xenopus
RT endoderm.";
RL Submitted (MAR-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY260732; AAP82289.1; -.
SQ SEQUENCE 462 AA; 53186 MW; 796F92774CC27721 CRC64;

Query Match 22.2%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 8.4e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

QY 10 PTKICVGCPRDPTNSPELEETLTHITIKLAENNAFYFKIDNVKKARQVQVQVAGK 65
DB 142 PGVILSTCP-DCTANEETPTITETADTLIAEYKNSNTTRYFKIDHIERVRSQWVGP 200
QY 66 KYFIDFVARETTCKESNEELTESC 90
DB 201 SYPIQFTIKETDCMKTQENWLSNC 225

RESULT 8
Q801E5 PRELIMINARY; PRT; 465 AA.
AC Q801E5
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical histidine-rich protein (Fragment).
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Pipidae;
OC Xenopodinae; Xenopus.
```

OX NCBI_TaxID=8355;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=22480013; PubMed=12591597;
 RA Chen Y., Jurgens K., Hollemann T., Claussen M., Ramadori G.,
 RA Pieler T.,
 RT "Cell-autonomous and signal-dependent expression of liver and
 RT intestine marker genes in pluripotent precursor cells from Xenopus
 RT embryos.";
 RL Mech. Dev. 120:277-288(2003).
 DR EMBL; AY186284; AAC31610.1; --
 DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; cystatin; 2.
 DR SMART; SM00043; CY; 2.
 KW Hypothetical protein.
 FT NON_TER 1
 SQ SEQUENCE 465 AA; 53528 MW; 0B403AB4P78BBFD4 CRC64;
 Query Match 22.2%; Score 152.5; DB 13; Length 465;
 Best Local Similarity 38.8%; Pred. No. 8.5e-07;
 Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;
 OX 10 PTKICVGPDRIPNTSPLEELTHT---ITKLAENNAFFYFKIDNVKARQVWVAGK 65
 Db 145 PGVILSTCP-DCPTANEETPTTETAEITLAEYNKDSNNTRYFKIDHIERVRSQWVGP 203
 OX 66 KVFIDFVARETTCKESNEELTESC 90
 Db 204 SYFIOTIKETDCMTQENVLSNC 228
 RESULT 9
 ID Q9EPX9 PRELIMINARY; PRT; 140 AA.
 AC Q9EPX9;
 DT 01-MAR-2001 (TrEMBLrel. 16, Created)
 DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE Cystatin C.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX STRAIN=BALB/c;
 RC MEDLINE=21010502; PubMed=11144350;
 RA Taupin P.J., Ray J., Fischer W.H., Suhr S.T., Hakansson K., Grubb A.,
 RA Gage F.H.;
 RT "FGF-2-Responsive neural stem cell proliferation requires Ccrg, a novel
 RT autocrine/paracrine cofactor.";
 RL Neuron 28:385-397(2000).
 DR EMBL; AF311741; AAG40283.1; --
 DR HSP; P01034; 1G96.
 DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; cystatin; 1.
 DR SMART; SM00043; CY; 1.
 DR PROSITE; PS00287; CYSTATIN; 1.
 FT CHAIN 21 140
 FT VARIANT 15 16 A -> G.
 FT VARIANT 84 84 L -> P.
 SQ SEQUENCE 140 AA; 15517 MW; 3A563406DD58D785 CRC64;
 Query Match 19.1%; Score 131.5; DB 11; Length 140;
 Best Local Similarity 27.8%; Pred. No. 2.6e-05;
 Matches 32; Conservative 26; Mismatches 48; Indels 9; Gaps 4;
 OX 15 VCCPRDIPNTSPLEELTHTITKLAENNAFFYFKIDNVKARQVWVAGKYYIDFVAR 74
 Db 30 LGAPEADANEGVRALDFAVSEYKNGSDNAYHSRAQVVRARKQLVAGVNYFLDVEMG 89

OX 75 ETTCSKESNEELTESC---ETKLGSLDCNAREVYVVPWEKKIYPTVTNNHCE 126
 Db 90 RITCTK-SQTNLTD-CFFHDQPHLMKALCSFQIYSPWK---GTHSLTNFSCK 138
 RESULT 10
 ID Q800S8 PRELIMINARY; PRT; 455 AA.
 AC Q800S8;
 DT 01-JUN-2003 (TrEMBLrel. 24, Created)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Fetuin-A.
 OS Brachydanio rerio (Zebrafish) (Danio rerio).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
 OC Cyprinidae; Danio.
 OX NCBI_TaxID=7955;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Liver;
 RA Jia F.;
 RT "Danio rerio fetuin-A.";
 RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AY217758; AAC61483.1; --
 DR GO; GO:0005874; C:microtubule; IEA.
 DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
 DR GO; GO:0005198; F:structural molecule activity; IEA.
 DR GO; GO:0007018; F:microtubule-based movement; IEA.
 DR InterPro; IPR002453; Beta_tubulin.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; cystatin; 1.
 DR SMART; SM00043; CY; 1.
 DR PROSITE; PS00228; TUBULIN B AUTOREG; 1.
 DR PROSITE; PS00228; TUBULIN B AUTOREG; 1.
 SQ SEQUENCE 455 AA; 50627 MW; D8228729268A2ACB CRC64;
 Query Match 18.0%; Score 123.5; DB 13; Length 455;
 Best Local Similarity 26.7%; Pred. No. 0.00063;
 Matches 32; Conservative 23; Mismatches 46; Indels 19; Gaps 4;
 OX 2 SGKDVQPTKICVGPDRIPNTSPLEELTHTITKLAENNAFFYFKIDNVKARQV- 60
 Db 134 SHEDLV---KKPCDGLPLHEPKALSVNAALAKFNKQSNHKSIFYKLMVGRISOW 189
 OX 61 VWAGKYYIDFVARETTCKESNEELTES------CETKLG-QSLDCNAREVY 106
 Db 190 MPMGQSYFTQFAIMETNCTKDAQNPQPEACKALCGDQATYFCCKSKVGSERPEVECIY 249
 RESULT 11
 ID Q8CB17 PRELIMINARY; PRT; 388 AA.
 AC Q8CB17;
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Fetuin beta.
 GN FETUB.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/60; TISSUE=Vagina;
 RX MEDLINE=22354683; PubMed=12466851;
 RA The FANTOM Consortium,
 RA the RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 RT 60,770 full-length cDNAs.";
 RL Nature 420:563-573(2002).
 RL EMBL; AK037043; BAC29682.1; --
 DR MGD; MGI:1890221; Fetub.

DT 01-OCT-2000 (TReMBLrel. 15, Created)
 DT 01-OCT-2000 (TReMBLrel. 15, Last sequence update)
 DT 01-JUN-2003 (TReMBLrel. 24, Last annotation update)
 DT 01-JUN-2003 (TReMBLrel. 24, Last annotation update)
 DE Ls-cystatin.
 OS Litomosoides sigmodontis.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
 OC Onchocercidae; Litomosoides.
 NCBI_TaxID=42156;
 [1]
 RN NCBI_TaxID=42156;
 RN
 RP SEQUENCE FROM N.A.
 RA Pfaff A.W., Hoffmann W.H., Taylor D.W., Schulz-Key H.;
 RT "Characterization and immunological properties of a cysteine protease
 RT inhibitor of the filarial parasite Litomosoides sigmodontis.";
 RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
 DR EMBL, AF229173; AAF35896.1; -;
 DR GO; GO:0004865; F:cysteine protease inhibitor activity; IEA.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; cystatin; 1.
 DR SMART; SM00043; Cy; 1.
 DR PROSITE; PS00287; CYSTATIN; 1.
 FT CHAIN 25 148
 FT SEQUENCE 148 AA; 16686 MW; 2950AA89CA5339C9 CRC64;
 SQ

 Query Match 16.5%; Score 113.5; DB 5; Length 148;
 Best Local Similarity 34.1%; Pred. No. 0.0017;
 Matches 30; Conservative 16; Mismatches 37; Indels 5; Gaps 3

 QY 27 ELEETLTHTITKLAENNAATEFYKIDNVKARQVWAGKYKVFIDFVARETTCCKESNEEL 86
 DB 49 EIQEMLPAILKVKQNSDNDALHLPKVLKVSQVWAGMYKKEIQVARSCKSSNEKI 108
 QY 87 -TESCETKLGOSLD--CNAEVYVPWE 111
 DB 109 DLKTC--KKLGHPDQIITLEVWEKAW 134

 RESULT 14
 Q80Y72
 ID Q80Y72 PRELIMINARY; PRT; 140 AA.
 AC Q80Y72;
 DT 01-JUN-2003 (TReMBLrel. 24, Created)
 DT 01-JUN-2003 (TReMBLrel. 24, Last sequence update)
 DT 01-OCT-2003 (TReMBLrel. 25, Last annotation update)
 DE Cystatin-like 1.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NCBI_TaxID=10090;
 [1]
 RN SEQUENCE FROM N.A.
 RP SEQUENCE FROM N.A.
 RA Scapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Vallalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Kettelman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
 RA Krzywinski M.I., Skalska U., Smallos D.E., Schnerch A., Schein J.E.,
 RA Jones S.J., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
 RN [2]

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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:04:32 ; Search time 8.636 Seconds
(without alignments)
765.738 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 697
Sequence: 1 GSKDFVQPTKICVGCPRD.....VPWEKKIYFTVTNNHCEPF 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_42:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	618	50.0	644	1 KNG HUMAN	P01042 homo sapien
2	440	64.0	436	1 KNL1_BOVIN	P01046 bos taurus
3	440	64.0	621	1 KXH1_BOVIN	P01044 bos taurus
4	413	60.1	434	1 KNL2_BOVIN	P01047 bos taurus
5	413	60.1	619	1 KXH2_BOVIN	P01045 bos taurus
6	413	60.1	661	1 KNG_MOUSE	O08677 mus musculus
7	410	59.7	639	1 KNG_RAT	P08934 rattus norv
8	388	56.5	430	1 KNT2_RAT	P08932 rattus norv
9	380	55.3	430	1 KNT1_RAT	P01048 rattus norv
10	171.5	25.0	144	1 CYTF_MOUSE	O09098 mus musculus
11	163.5	23.8	145	1 CYTF_HUMAN	O76096 homo sapien
12	138.5	20.2	146	1 CYTC_MACMU	O19092 macaca mula
13	138.5	20.2	149	1 CYTM_HUMAN	Q15828 homo sapien
14	137.5	20.0	148	1 CYTC_BOVIN	P01035 bos taurus
15	135	19.7	378	1 FETB_RAT	Q09x79 rattus norv
16	132.5	19.3	146	1 CYTC_HUMAN	P01034 homo sapien
17	132.5	19.3	146	1 CYTC_SALIC	O19093 salmire sci
18	131.5	19.1	382	1 FETB_HUMAN	Q9UGM5 homo sapien
19	130	18.9	127	1 CYTC_RAT	P14841 rattus norv
20	129.5	18.9	140	1 CYTC_MOUSE	P21460 mus musculus
21	128	18.6	111	1 CYT_BITAR	P08935 bitis ariet
22	124.5	18.1	141	1 CYTT_HUMAN	P05228 homo sapien
23	124.5	18.1	148	1 CYTC_RABIT	O97862 oryctolagus
24	122.5	17.8	116	1 CYT_COTJA	P81061 coturnix co
25	119	17.3	388	1 FETB_MOUSE	Q9GXC1 mus musculus
26	118.5	17.2	139	1 CYT_CHICK	P01038 gallus gall
27	113	16.4	141	1 CYTS_RAT	P19313 rattus norv
28	109.5	15.9	141	1 CYTN_HUMAN	P01037 homo sapien
29	108.5	15.8	141	1 CYTS_HUMAN	P01036 homo sapien
30	107	15.6	130	1 CYT_ONCKE	Q98967 oncorhynch
31	105.5	15.4	162	1 CYTX_ONCVO	P22085 onchocerca
32	105	15.3	130	1 CYT_ONCMY	Q91195 oncorhynch
33	104	15.1	129	1 CYT_CYPCA	P35481 cyprinus ca

ALIGNMENTS

RESULT 1

ID	KNG_HUMAN	STANDARD;	PRT;	644 AA.
AC	P01042; P01043;			
DT	21-JUL-1986 (Rel. 01, Created)			
DT	01-FEB-1996 (Rel. 33, Last sequence update)			
DT	10-OCT-2003 (Rel. 42, Last annotation update)			
DE	Kinogen precursor (Alpha-2-thiol proteinase inhibitor) [Contains: Bradykinin].			
GN	KNG.			
OS	Homo sapiens (Human).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
OX	NCBI_TaxID=9606;			
RN	[1]			
RP	SEQUENCE FROM N.A. (ISOFORMS HMW AND LMW).			
RC	TISSUE=Liver;			
RX	MEDLINE=85234582; PubMed=2989293;			
RA	Takagaki Y., Kitamura N., Nakanishi S.;			
RT	"Cloning and sequence analysis of cDNAs for human high molecular weight and low molecular weight prekininogens. Primary structures of two human prekininogens.";			
RT	J. Biol. Chem. 260:8601-8609(1985).			
RN	[2]			
RP	GENE STRUCTURE			
RX	MEDLINE=85234583; PubMed=2989294;			
RA	Kitamura N., Kitagawa H., Fukushima D., Takagaki Y., Miyata T., Nakanishi S.;			
RT	"Structural organization of the human kininogen gene and a model for its evolution.";			
RT	J. Biol. Chem. 260:8610-8617(1985).			
RN	[3]			
RP	SEQUENCE OF 1-401 FROM N.A.			
RX	MEDLINE=85122621; PubMed=6441591;			
RA	Okubo I., Kurachi K., Takasawa T., Shiokawa H., Sasaki M.;			
RT	"Isolation of a human cDNA for alpha 2-thiol proteinase inhibitor and its identity with low molecular weight kininogen.";			
RT	Biochemistry 23:5691-5697(1984).			
RN	[4]			
RP	SEQUENCE OF 379-644.			
RX	MEDLINE=86030270; PubMed=4054110;			
RA	Lottspeich F., Kellermann J., Henschel A., Foertsch B., Mueller-Esterl W.;			
RT	"The amino acid sequence of the light chain of human high-molecular-mass kininogen.";			
RT	Eur. J. Biochem. 152:307-314(1985).			
RN	[5]			
RP	SEQUENCE OF 381-389.			
RX	MEDLINE=90255622; PubMed=4952632;			
RA	Pierce J.V.;			
RT	"Structural features of plasma kinins and kininogens.";			
RT	Fed. Proc. 27:52-57(1968).			
RN	[6]			
RP	DISULFIDE BONDS.			
RA	Sueyoshi T., Miyata T., Kato H., Iwanaga S.;			
RT	"Disulfide bonds in bovine HMW kininogens.";			

P31727 sarcophaga
Q94269 mus musculus
Q9h114 homo sapien
P27666 mus musculus
P29699 mus musculus
O60576 homo sapien
P81714 najia atra (P81714)
Q9h112 homo sapien
P28325 homo sapien
P31726 zea mays (m
P24090 rattus norv
P02765 homo sapien

34 102 14.8 122 1 CYTA_SARPE
35 101 14.7 139 1 CS11_MOUSE
36 95 13.8 165 1 CSTL_HUMAN
37 94.5 13.8 142 1 CST8_MOUSE
38 94.5 13.8 345 1 A2HS_MOUSE
39 94 13.7 142 1 CST8_HUMAN
40 93 13.5 99 1 CYT_NAJAT
41 93 13.5 137 1 CS11_HUMAN
42 93 13.5 142 1 CYTD_HUMAN
43 91.5 13.3 135 1 CYTI_NAJZE
44 91.5 13.3 352 1 A2HS_RAT
45 88 12.8 367 1 A2HS_HUMAN

Seikagaku 56:808-808(1984).
[7]
CARBOHYDRATE-LINKAGE SITE ASN-294.
MEDLINE-22660472; PubMed-12754519;
Zhang H., Li X.-J., Martin D.B., Abersold R.;
Identification and quantification of N-linked glycoproteins using
hydrazide chemistry, stable isotope labeling and mass spectrometry.";
Nat. Biotechnol. 21:660-666(2003).
-!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
HMM-kininogen plays an important role in blood coagulation by
helping to position optimally prekallikrein and factor XI next to
factor XII; (3) HMM-kininogen inhibits the thrombin- and plasmin-
induced aggregation of thrombocytes; (4) the active peptide
bradykinin that is released from HMM-kininogen shows a variety of
physiological effects: (4A) influence in smooth muscle
contraction, (4B) induction of hypotension, (4C) natriuresis and
diuresis, (4D) decrease in blood glucose level, (4E) it is a
mediator of inflammation and causes (4E1) increase in vascular
permeability, (4E2) stimulation of nociceptors (4E3) release of
other mediators of inflammation (e.g. prostaglandins), (4F) it has
a cardioprotective effect (directly via bradykinin action,
indirectly via endothelium-derived relaxing factor action); (5)
LMM-kininogen inhibits the aggregation of thrombocytes; (6) LMM-
kininogen is in contrast to HMM-kininogen not involved in blood
clotting.
-!- SUBCELLULAR LOCATION: Secreted.
-!- ALTERNATIVE PRODUCTS:
Event=Alternative splicing; Named isoforms=2;
Name=HMM;
IsoId=P01042-1; Sequence=Displayed;
Name=LMM;
IsoId=P01042-2; Sequence=VSP_001261, VSP_001262;
-!- TISSUE SPECIFICITY: Plasma.
-!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
-!- SIMILARITY: Contains 3 cystatin-like domains.

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DR EMBL; K02566; AAB35497.1; --
DR EMBL; M11437; AAB59550.1; --
DR EMBL; M11438; AAB59550.1; JOINED.
DR EMBL; M11521; AAB59550.1; JOINED.
DR EMBL; M11522; AAB59550.1; JOINED.
DR EMBL; M11523; AAB59550.1; JOINED.
DR EMBL; M11524; AAB59550.1; JOINED.
DR EMBL; M11525; AAB59550.1; JOINED.
DR EMBL; M11526; AAB59550.1; JOINED.
DR EMBL; M11527; AAB59550.1; JOINED.
DR EMBL; M11528; AAB59550.1; JOINED.
DR EMBL; M11437; AAB59551.1; --
DR EMBL; M11438; AAB59551.1; JOINED.
DR EMBL; M11521; AAB59551.1; JOINED.
DR EMBL; M11522; AAB59551.1; JOINED.
DR EMBL; M11523; AAB59551.1; JOINED.
DR EMBL; M11524; AAB59551.1; JOINED.
DR EMBL; M11525; AAB59551.1; JOINED.
DR EMBL; M11526; AAB59551.1; JOINED.
DR EMBL; M11527; AAB59551.1; JOINED.
DR EMBL; M11528; AAB59551.1; JOINED.
DR PIR; A01278; KGHUHL.
DR PIR; A01280; KGHUHL.
DR SWISS-2DPAGE; P01042; HUMAN.
DR Genew; HGNC:6383; KNG.
DR MIM; 228960; --
DR GO; GO:0007596; P:blood coagulation; NAS.
DR GO; GO:0030146; P:diuresis; NAS.
DR GO; GO:0006954; P:inflammatory response; NAS.

DR GO; GO:0030147; P:natriuresis; NAS.
DR GO; GO:0006939; P:smooth muscle contraction; NAS.
DR InterPro; IPR000010; Cystatin.
DR InterPro; IPR002395; Kininogen.
DR Pfam; PF00031; cystatin; 3.
DR PRINTS; PR00334; KININOGEN.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSSTATIN; 2.
KW Glycopoltein; Plasma; Repeat; Thiol protease inhibitor; Vasodilator;
KW Bradykinin; Blood coagulation; Inflammatory response; Signal;
KW Alternative splicing; Pyrrolidone carboxylic acid.
FT SIGNAL 1 18
FT CHAIN 19 644 KININOGEN.
FT CHAIN 19 380 KININOGEN HEAVY CHAIN.
FT PEPTIDE 381 389 BRADYKININ.
FT CHAIN 390 644 KININOGEN LIGHT CHAIN.
FT DOMAIN 19 136 CYSTATIN-LIKE 1.
FT DOMAIN 137 258 CYSTATIN-LIKE 2.
FT DOMAIN 259 380 CYSTATIN-LIKE 3.
FT DOMAIN 420 510 HIS-RICH
(ASSOCIATED WITH CLOTTING ACTIVITY).
FT REPEAT 420 449
FT REPEAT 450 479
FT REPEAT 480 510
FT MOD_RES 19 19
FT DISULFID 28 614
FT DISULFID 83 94
FT DISULFID 107 126
FT DISULFID 142 145
FT DISULFID 206 218
FT DISULFID 229 248
FT DISULFID 264 267
FT DISULFID 328 340
FT DISULFID 351 370
FT CARBOHYD 48 48 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 169 169 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 205 205 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 294 294 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 401 401 O-LINKED.
FT CARBOHYD 533 533 O-LINKED.
FT CARBOHYD 542 542 O-LINKED.
FT CARBOHYD 546 546 O-LINKED.
FT CARBOHYD 557 557 O-LINKED.
FT CARBOHYD 571 571 O-LINKED.
FT CARBOHYD 577 577 O-LINKED.
FT CARBOHYD 593 593 O-LINKED.
FT CARBOHYD 628 628 O-LINKED.
FT VARSPLIC 402 427
FT VARSPLIC 428 644
FT VARSPLIC 593 593
FT CONFLICT 593 593
FT CONFLICT 644 644
SQ SEQUENCE 644 AA; 71945 MW; 3132B4CBAF8FB7E CRC64;
Query Match 90.0%; Score 618; DB 1; Length 644;
Best Local Similarity 100.0%; Pred. No. 3.4e-51;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTITITKLAENNATFYFKIDNVKARQV 62
DB 253 GKDFVQPTKICVGCPRDIPNTSPLEETLTITITKLAENNATFYFKIDNVKARQV 312
QY 63 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKITYPTV 118
DB 313 AGKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKITYPTV 368
RESULT 2
KNLI BOVIN STANDARD; PRT; 436 AA.
ID KNLI BOVIN
AC P01046;
DT 21-JUL-1986 (Rel. 01, Created)

DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Kininogen, LMW I precursor (Thiol proteinase inhibitor) [Contains:
 DE Bradykinin].
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OC NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=93117859; PubMed=6572010;
 RA Nawa H., Kitamura N., Hirose T., Asai M., Inayama S., Nakanishi S.;
 RT "Primary structures of bovine liver low molecular weight kininogen
 RT precursors and their two mRNAs.";
 RL Proc. Natl. Acad. Sci. U.S.A. 80:190-94 (1983).
 RN [2]
 RP SEQUENCE OF 19-378.
 RX MEDLINE=87137530; PubMed=3546295;
 RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 RA Miyata T., Iwanaga S.;
 RT "Bovine high molecular weight kininogen. The amino acid sequence,
 RT positions of carbohydrate chains and disulfide bridges in the heavy
 RT chain portion.";
 RL J. Biol. Chem. 262:2768-2779 (1987).
 CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 CC LMW-kininogen inhibits the aggregation of thrombocytes; (3) the
 CC active peptide kallidin that is released from LMW-kininogen shows
 CC a variety of physiological effects: (3A) influence in smooth
 CC muscle contraction, (3B) induction of hypotension, (3C)
 CC natriuresis and diuresis (kidney).
 CC -!- SUBCELLULAR LOCATION: Extracellular.
 CC -!- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=LMW I;
 CC IsoId=P01046-1; Sequence=Displayed;
 CC Name=HMW I;
 CC IsoId=P01044-1; Sequence=External;
 CC -!- TISSUE SPECIFICITY: Plasma.
 CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -!- MISCELLANEOUS: LMW-kininogen is in contrast to HMW-kininogen not
 CC involved in blood clotting.
 CC -!- SIMILARITY: Contains 3 cystatin-like domains.
 CC
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 CC
 CC EMBL; V00426; CAA23709.1; ..
 DR PIR; A01283; KGBOL1.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; cystatin; 3.
 DR SMART; SM00043; CY; 3.
 DR PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
 KW Thiol protease inhibitor; Bradykinin; Signal;
 KW Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 436 KININOGEN, LMW I.
 FT CHAIN 19 378 HEAVY CHAIN.
 FT PEPTIDE 380 388 BRADYKININ.
 FT CHAIN 389 436 LIGHT CHAIN.
 FT DOMAIN 19 135 CYSTATIN-LIKE 1.
 FT DOMAIN 136 257 CYSTATIN-LIKE 2.
 FT DOMAIN 258 378 CYSTATIN-LIKE 3.
 FT MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
 FT MOD RES 57 87 N-LINKED (GLCNAC. . .).
 FT CARBOHYD 136 136 O-LINKED (PARTIAL).
 FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
 FT

FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
 FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
 FT DISULFID 27 406 INTERCHAIN.
 FT DISULFID 82 93
 FT DISULFID 106 125
 FT DISULFID 141 144
 FT DISULFID 205 217
 FT DISULFID 228 247
 FT DISULFID 263 266
 FT DISULFID 327 339
 FT DISULFID 350 369
 FT CONFLICT 295 295
 SQ SEQUENCE 436 AA: 48427 MW: 48427 MW: F01F7EB6914BCE6C CRC64;
 A -> T (IN REF. 1; CAA23709).
 Query Match 64.0%; Score 440; DB 1; Length 436;
 Best Local Similarity 70.4%; Pred. No. 1.8e-34;
 Matches 81; Conservative 14; Mismatches 20; Indels 0; Gaps 0;
 QY 4 KDFVQPPKICVCGPRDPTSPSELEBEVLTHTITKLNENNAFFYFKIDNVKKAQVQVA 63
 DB 253 KDFVQPPKICVCGPRDPTSPSELEBEVLTHTITKLNENNAFFYFKIDNVKKAQVQVA 312
 QY 64 GKYYFDVARETTCKSKESNEELTESCTKLGQSLDCNAEYVYVWPEKKIYPTV 118
 DB 313 GLKYSIVFIARETTCKSKESNEELTKSCENITHGQILHCDANVYVWPEKKIYPTV 367
 RESULT 3
 KNHL BOVIN STANDARD; PRT; 621 AA.
 ID KNHL_BOVIN
 AC P01044;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Kininogen, HMW I precursor (Thiol proteinase inhibitor) [Contains:
 DE Bradykinin].
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OC NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=84014106; PubMed=6571699;
 RA Kitamura N., Takagaki Y., Furuto S., Tanaka T., Nawa H., Nakanishi S.;
 RT "A single gene for bovine high molecular weight and low molecular
 RT weight kininogens.";
 RL Nature 305:545-549 (1983).
 RL [2]
 RP SEQUENCE OF 19-378.
 RX MEDLINE=87137530; PubMed=3546295;
 RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 RA Miyata T., Iwanaga S.;
 RT "Bovine high molecular weight kininogen. The amino acid sequence,
 RT positions of carbohydrate chains and disulfide bridges in the heavy
 RT chain portion.";
 RL J. Biol. Chem. 262:2768-2779 (1987).
 RN [3]
 RP SEQUENCE OF 378-393.
 RX MEDLINE=70180420; PubMed=4986212;
 RA Kato H., Nagasawa S., Suzuki T.;
 RT "Studies on the structure of bovine kininogen: cleavages of disulfide
 RT bonds and of methionyl bonds in kininogen-II.";
 RL J. Biochem. 67:313-323 (1970).
 RN [4]
 RP SEQUENCE OF 458-498.
 RX MEDLINE=75170265; PubMed=1169237;
 RA Han Y.N., Komiya M., Iwanaga S., Suzuki T.;
 RT "Studies on the primary structure of bovine high-molecular-weight
 RT kininogen. Amino acid sequence of a fragment ('histidine-rich
 RT peptide') released by plasma kallikrein.";
 RL J. Biochem. 77:155-68 (1975).
 CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)

HMW-kininogen plays an important role in blood coagulation by helping to position optimally prekallikrein and factor XI next to factor XII. (3) HMW-kininogen inhibits the thrombin- and plasmin-induced aggregation of thrombocytes; (4) the active peptide bradykinin that is released from HMW-kininogen shows a variety of physiological effects: (4A) influence in smooth muscle contraction, (4B) induction of hypotension, (4C) natriuresis and diuresis, (4D) decrease in blood glucose level. (4E) it is a mediator of inflammation and causes (4E1) increase in vascular permeability, (4E2) stimulation of nociceptors (4E3) release of other mediators of inflammation (e.g. prostaglandins), (4F) it has a cardioprotective effect (directly via bradykinin action, indirectly via endothelium-derived relaxing factor action).

-1- SUBCELLULAR LOCATION: Extracellular.

-1- ALTERNATIVE PRODUCTS:

Event=Alternative splicing; Named isoforms=2;

Name=HMW II;

isoId=P01044-1; Sequence=Displayed;

Name=LMW I;

isoId=P01046-1; Sequence=External;

-1- TISSUE SPECIFICITY: Plasma.

-1- PTM: Bradykinin is released from kininogen by plasma kallikrein.

-1- SIMILARITY: Contains 3 cystatin-like domains.

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EMBL; V01491; CAA24735.1; -

PIR; A01281; KGBOL1.

InterPro; IPR000010; Cystatin.

InterPro; IPR002395; Kininogen.

Pfam; PF00031; Cystatin; 3.

PRINTS; PR00334; KININOGEN.

SMART; SM00043; CY; 3.

PROSITE; PS00287; CYSTATIN; 2.

KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;

KW Thiol protease inhibitor; Bradykinin; Blood coagulation;

KW Inflammatory response; Signal; Pyrrolidone carboxylic acid.

FT SIGNAL 1 18 PROBABLE

FT CHAIN 19 621 KININOGEN, HMW I.

FT CHAIN 19 378 HEAVY CHAIN.

FT PEPTIDE 380 388 BRADYKININ.

FT CHAIN 389 621 LIGHT CHAIN.

FT DOMAIN 19 135 CYSTATIN-LIKE 1.

FT DOMAIN 136 257 CYSTATIN-LIKE 2.

FT DOMAIN 258 378 CYSTATIN-LIKE 3.

FT MOD_RES 19 19 PYRROLIDONE CARBOXYLIC ACID.

FT CARBOHYD 87 87 N-LINKED (GLCNAC. . .).

FT CARBOHYD 136 136 O-LINKED (PARTIAL. . .).

FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).

FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).

FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).

FT DISULFID 27 591 INTERCHAIN.

FT DISULFID 82 93

FT DISULFID 106 125

FT DISULFID 141 144

FT DISULFID 205 217

FT DISULFID 228 247

FT DISULFID 263 266

FT DISULFID 327 339

FT DISULFID 350 369

FT SEQUENCE 621 AA; 68890 MW; D16850BEFE3C55CD CRC64;

Query Match 64.0%; Score 440; DB 1; Length 621;

Best Local Similarity 70.4%; Pred. No. 2.7e-34;

Matches 81; Conservative 14; Mismatches 20; Indels 0; Gaps 0;

Db 253 KDFVQPTRLCAGCPKIPVDSDEPLSHSLAKLNHEDGAFYFKIDIVKATQVVA 312

QY 64 GKXYFIDFVARETTCSKESNEELTESCETKKLGSLDCNAEVYVWPWEKIIPTV 118

Db 313 GLKYSIVFIARETTCSKSGSNEELTKSCEINIHGQILHCDANVYVWPWEKIVPTV 367

RESULT 4

KML2_BOVIN STANDARD; PRT; 434 AA.

ID KML2_BOVIN AC P01047;

DT 21-JUL-1986 (Rel. 01, Created)

DT 21-JUL-1986 (Rel. 01, Last sequence update)

DT 15-MAR-2004 (Rel. 43, Last annotation update)

DE Kininogen, LMW II precursor (Thiol proteinase inhibitor) [Contains: Bradykinin]

OS Bos taurus (Bovine).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovidae; Bovinae; Bos.

OX NCBI_TaxID=9913;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=83117859; PubMed=6572010;

RA Nawa H., Kitamura N., Hirose T., Asai M., Inayama S., Nakarishi S.; Miyata T., Iwanaga S.;

RT "Primary structures of bovine liver low molecular weight kininogen precursors and their two mRNAs";

RL Proc. Natl. Acad. Sci. U.S.A. 80:90-94 (1983).

RN [2]

RP SEQUENCE OF 19-376.

RX MEDLINE=87137530; PubMed=3546295;

RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H., Miyata T., Iwanaga S.;

RT "Bovine high molecular weight kininogen. The amino acid sequence, positions of carbohydrate chains and disulfide bridges in the heavy chain portion.";

RL J. Biol. Chem. 262:2768-2779 (1987).

CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2) active peptide kallidin that is released from LMW-kininogen shows a variety of physiological effects: (3A) influence in smooth muscle contraction, (3B) induction of hypotension, (3C) natriuresis and diuresis (kidney).

CC -1- SUBCELLULAR LOCATION: Extracellular.

CC -1- ALTERNATIVE PRODUCTS:

Event=Alternative splicing; Named isoforms=2;

Name=LMW II;

isoId=P01047-1; Sequence=Displayed;

Name=HMW II;

isoId=P01045-1; Sequence=External;

-1- TISSUE SPECIFICITY: Plasma.

-1- PTM: Bradykinin is released from kininogen by plasma kallikrein.

-1- MISCELLANEOUS: LMW-kininogen is in contrast to HMW-kininogen not involved in blood clotting.

-1- SIMILARITY: Contains 3 cystatin-like domains.

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EMBL; V00427; CAA23710.1; -

PIR; A01284; KGBOL2.

HSSP; P01038; 1A90.

InterPro; IPR000010; Cystatin.

Pfam; PF00031; Cystatin; 3.

SMART; SM00043; CY; 3.

PROSITE; PS00287; CYSTATIN; 2.

KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;

```
KW Thiol protease inhibitor; Bradykinin; Signal;
KW Pyrrolidone carboxylic acid.
FT SIGNAL 1 18
FT CHAIN 19 434 KININOGEN, LMW II.
FT CHAIN 19 376 HEAVY CHAIN.
FT PEPTIDE 378 386 BRADYKININ.
FT CHAIN 387 434 LIGHT CHAIN.
FT CHAIN 387 434 CYSSTATIN-LIKE 1.
FT DOMAIN 136 256 CYSSTATIN-LIKE 2.
FT DOMAIN 136 256 CYSSTATIN-LIKE 3.
FT MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
FT CARBOHYD 87 87 N-LINKED (GLCNAC. . .).
FT CARBOHYD 136 136 O-LINKED (PARTIAL. . .).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
FT CARBOHYD 280 280 N-LINKED (GLCNAC. . .).
FT DISULFID 27 404 INTERCHAIN.
FT DISULFID 82 93
FT DISULFID 106 125
FT DISULFID 141 144
FT DISULFID 205 217
FT DISULFID 228 247
FT DISULFID 261 264
FT DISULFID 325 337
FT DISULFID 348 367
SQ SEQUENCE 434 AA; 48148 MW; 73A7079DE3E03430 CRC64;

Query Match 60.1%; Score 413; DB 1; Length 434;
Best Local Similarity 67.2%; Pred. No. 6.6e-32;
Matches 78; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

QY 3 GKDFVQPTKLCVGPDPINSPSELETLTHITKLNAENNAFYFKIDNVKKARQVQV 62
D 252 GEDFL--PPWVCVCPKPIPVDSPLDEALNHSIAKLNAEHDGTFYKIDTVKATQVQV 309
QY 63 AGKYFIDFVARETTCKSENEELTESCETKLGQSLDCNAEVVYVPEKKIYPTV 118
D 310 GGLKYSIVFIARETTCKSGSNEELTKSCEINHGQILHCDANVYVPEEKVYPTV 365

RESULT 5
ID KXH2_BOVIN STANDARD; PRT; 619 AA.
AC P01045;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Kininogen, HMW II precursor (thiol proteinase inhibitor) [Contains:
OS Bradykinin].
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxId=9913;
RN [1]
RN SEQUENCE FROM N.A.
RX MEDLINE=84014106; PubMed=6571699;
RA Kitamura N., Takagaki Y., Furuto S., Tanaka T., Nawa H., Nakanishi S.;
RT "A single gene for bovine high molecular weight and low molecular
RT weight kininogens."
RL Nature 305:545-549 (1983).
RN [2]
RN SEQUENCE OF 19-376.
RX MEDLINE=87137530; PubMed=3546295;
RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
RA Miyata T., Iwanaga S.;
RT "Bovine high molecular weight kininogen. The amino acid sequence,
RT positions of carbohydrate chains and disulfide bridges in the heavy
RT chain portion."
RL J. Biol. Chem. 262:2768-2779 (1987).
RN [3]
RN SEQUENCE OF 376-391.
RX MEDLINE=70180420; PubMed=4986212;
Kato H., Nagasawa S., Suzuki T.;
RT "Studies on the structure of bovine kininogen: cleavages of disulfide
RT bonds and of methionyl bonds in kininogen-II."
J. Biochem. 67:313-323 (1970).
RN [4]
RN SEQUENCE OF 387-455.
RX MEDLINE=76260155; PubMed=956151;
Han Y.N., Kato H., Iwanaga S., Suzuki T.;
RT "Primary structure of bovine plasma high-molecular-weight kininogen.
RT The amino acid sequence of a glycopeptide portion (fragment 1)
RT following the C-terminus of the bradykinin moiety."
J. Biochem. 79:1201-1222 (1976).
RN [5]
RN SEQUENCE OF 456-496.
RX MEDLINE=75170265; PubMed=1169237;
Han Y.N., Komiyama M., Iwanaga S., Suzuki T.;
RT "Studies on the primary structure of bovine high-molecular-weight
RT kininogen. Amino acid sequence of a fragment ('histidine-rich
RT peptide') released by plasma kallikrein."
J. Biochem. 77:55-68 (1975).
CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
HMW-kininogen plays an important role in blood coagulation by
helping to position optimally prekallikrein and factor XI next to
factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
induced aggregation of thrombocytes; (4) the active peptide
bradykinin that is released from HMW-kininogen shows a variety of
physiological effects: (4A) influence in smooth muscle
contraction, (4B) induction of hypotension, (4C) natriuresis and
diuresis, (4D) decrease in blood glucose level, (4E) it is a
mediator of inflammation and causes (4E1) increase in vascular
permeability, (4E2) stimulation of nociceptors (4E3) release of
other mediators of inflammation (e.g. prostaglandins), (4F) it has
a cardioprotective effect (directly via bradykinin action,
indirectly via endothelium-derived relaxing factor action).
CC -1- SUBCELLULAR LOCATION: Extracellular.
CC -1- ALTERNATIVE PRODUCTS:
Event=Alternative splicing; Named isoforms=2;
Name=HMW II;
IsoId=P01045-1; Sequence=Displayed;
Name=LMW II;
IsoId=P01047-1; Sequence=External;
CC -1- TISSUE SPECIFICITY: Plasma.
CC -1- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -1- SIMILARITY: Contains 3 cystatin-like domains.
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entities requires a license agreement (See http://www.isb-sib.ch/announce/
or send an email to licenses@isb-sib.ch).
CC EMBL: V01492; CAA24736.1; -.
DR PIR: A01282; KGB0H2.
DR HSP: P01038; IAS0.
DR InterPro: IPR000010; Cystatin.
DR InterPro: IPR002395; Kininogen.
DR Pfam: PF00031; Cystatin; 3.
DR PRINTS: PR00334; KININOGEN.
DR SMART: SM00043; CY. 3.
DR PROSITE: PS00287; CYSSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
KW Thiol protease inhibitor; Bradykinin; Blood coagulation; Signal;
KW Inflammatory response; Pyrrolidone carboxylic acid.
FT SIGNAL 1 18 KININOGEN, HMW II.
FT CHAIN 19 619 HEAVY CHAIN.
FT CHAIN 19 376 BRADYKININ.
FT PEPTIDE 378 386 LIGHT CHAIN.
FT CHAIN 387 619 CYSSTATIN-LIKE 1.
FT DOMAIN 19 135 CYSSTATIN-LIKE 2.
FT DOMAIN 136 256
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FT DOMAIN 257 376 CYPSTATIN-LIKE 3.
FT MOD_RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
FT CARBOHYD 87 87 N-LINKED (GLCNAC. . .)
FT CARBOHYD 136 136 O-LINKED (PARTIAL. . .)
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL. . .)
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .)
FT CARBOHYD 280 280 N-LINKED (GLCNAC. . .)
FT CARBOHYD 400 400 O-LINKED.
FT DISULFID 27 589 INTERCHAIN.
FT DISULFID 82 93
FT DISULFID 106 125
FT DISULFID 141 144
FT DISULFID 205 217
FT DISULFID 228 247
FT DISULFID 261 264
FT DISULFID 325 337
FT DISULFID 348 367
FT VARIANT 398 398 T -> P.
FT VARIANT 401 401 L -> V.
FT VARIANT 454 454 H -> K.
SQ SEQUENCE 619 AA; 68710 MW; F04320A8EB0E0DA CRC64;

Query Match
Best Local Similarity 60.1%; Score 413; DB 1; Length 619;
Matches 78; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

QY 3 GKDFVPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARVQV 62
Db 252 GEDFL--PPMVCVCGPKPVPDPSDLAEALNSIAKNAEHDTGYFKIDTVKKATQV 309
QY 63 AKKYPIDVARETTCKESNELTESCTKLGSLDCAEYVYVPEKKIYPTV 118
Db 310 GGLKYSIVFIARETTCKSGNELTKSCBEINTHGQILCHDANVYVPEEKYPTV 365

RESULT 6
KING_MOUSE STANDARD; PRT; 661 AA.
AC O08677; O08676; Q91XK5;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Kininogen precursor [Contains: Bradykinin].
GN KING.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORMS HMW AND LMW).
RC STRAIN=C57BL/6 X CBA; TISSUE=Liver;
RX MEDLINE=97342556; PubMed=9199253;
RA Okazaki Y., Furuno M., Kasukawa T., Adachi J., Bono H., Kondo S.,
RA Nikaide I., Osato N., Saito R., Suzuki H., Yamanaoka I., Kiyosawa H.,
RA Yagi K., Tomaru Y., Hasegawa Y., Nogami A., Schonbach C., Gojobori T.,
RA Baldarelli R., Hill D.P., Bult C., Hume D.A., Quackenbush J.,
RA Schriml L.M., Khanpin A., Matsuda H., Batalov S., Beisel K.W.,
RA Blake J.A., Bradt D., Brusio V., Chothia C., Corbani L.E., Cousins S.,
RA Dalla E., Dragani T.A., Fletcher C.F., Forrest A., Frazer K.S.,
RA Gaasterland T., Gariboldi M., Gissi C., Godzik A., Gough J.,
RA Grimmond S., Gustincich S., Hirokawa N., Jackson I.J., Jarvis E.D.,
RA Kanai A., Kawaji H., Kawasawa Y., Kedzierski R.M., King B.L.,
RA Konagaya A., Kurochkin I.V., Lee Y., Lenhard B., Lyons P.A.,
RA Maglott D.R., Maltais L., Marchionni L., McKenzie L., Miki H.,

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RA Nagashima T., Numata K., Okido T., Pavan W.J., Pertea G., Pesole G.,
RA Petrovsky N., Pillai R., Pontius J.U., Qi D., Ramachandran S.,
RA Ravasi T., Reed J.C., Reed D.J., Reid J., Ring B.Z., Ringwald M.,
RA Sardinelli A., Schneider C., Sempie C.A., Setou M., Shimada K.,
RA Sultana R., Takenaka Y., Taylor M.S., Teasdale R.D., Tomita M.,
RA Verardo R., Wagner L., Wahlestedt C., Wang Y., Watanabe Y., Wells C.,
RA Wilmig L.G., Wynshaw-Boris A., Yanagisawa M., Yang I., Yang L.,
RA Yuan Z., Zavolan M., Zhu Y., Zimmer A., Carninci P., Hayatsu N.,
RA Hirozane-Kishikawa T., Konno H., Nakamura M., Sakazume N., Sato K.,
RA Shiraki T., Waki K., Kawai J., Aizawa K., Arakawa T., Fukuda S.,
RA Hara A., Hashizume W., Imotani K., Ishii Y., Itoh M., Kagawa I.,
RA Miyazaki A., Sakai K., Sasaki D., Shibata K., Shinagawa A.,
RA Yasunishi A., Yoshino M., Waterston R., Lander E.S., Rogers J.,
RA Birney E., Hayashizaki Y.;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs.";
RL Nature 420:563-573 (2002).
RN [3]
RP SEQUENCE FROM N.A. (ISOFORM LMW).
RC TISSUE=Liver;
RX MEDLINE=22388257; PubMed=12477932;
RA Srausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wegner L., Shennen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buettow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.F., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Besak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettaman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting N., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalios D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC HMW-kininogen plays an important role in blood coagulation by
CC helping to position optimally prekallikrein and factor XI next to
CC induced aggregation of thrombocytes; (4) the active peptide
CC bradykinin that is released from HMW-kininogen shows a variety of
CC physiological effects: (4A) influence in smooth muscle
CC contraction, (4B) induction of hypotension, (4C) natriuresis and
CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
CC mediator of inflammation and causes (4E1) increase in vascular
CC permeability, (4E2) stimulation of nociceptors (4E3) release of
CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action); (5)
CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
CC kininogen is in contrast to HMW-kininogen not involved in blood
CC clotting (By similarity).
CC -1- SUBCELLULAR LOCATION: Secreted.
CC -1- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=HMW;
CC IsoId=O08677-1; Sequence=Displayed;
CC Name=LMW;
CC IsoId=O08677-2; Sequence=VSP_001263, VSP_001264;
CC -1- TISSUE SPECIFICITY: Plasma.
CC -1- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -1- SIMILARITY: Contains 3 cystatin-like domains.
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DR EMBL; M11884; AAA41487.1; -
DR EMBL; M14369; AAA41484.1; -
DR EMBL; M14369; AAA41485.1; ALT_SEQ.
DR EMBL; M16455; AAA41482.1; -
DR PIR; A25486; A25486.
DR PIR; A28055; A28055.
DR InterPro; IPR000010; Cystatin.
DR PRINTS; PR00031; cystatin; 3.
DR SMART; SM00334; KININOGEN.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Thiol protease inhibitor; Vasodilator;
KW Bradykinin; Blood coagulation; Inflammatory response; Signal;
KW Alternative splicing; Multigene family.
FT SIGNAL 1 18
FT CHAIN 19 639 KININOGEN.
FT CHAIN 19 380 KININOGEN HEAVY CHAIN.
FT PEPTIDE 380 389 BRADYKININ.
FT CHAIN 391 639 KININOGEN LIGHT CHAIN.
FT DOMAIN 19 136 CYSTATIN-LIKE 1.
FT DOMAIN 137 288 CYSTATIN-LIKE 2.
FT DOMAIN 259 380 CYSTATIN-LIKE 3.
FT DOMAIN 439 514 HIS-RICH.
FT DISULFID 28 609 INTERCHAIN (BY SIMILARITY).
FT DISULFID 83 94 BY SIMILARITY.
FT DISULFID 107 126 BY SIMILARITY.
FT DISULFID 142 145 BY SIMILARITY.
FT DISULFID 206 218 BY SIMILARITY.
FT DISULFID 229 248 BY SIMILARITY.
FT DISULFID 264 267 BY SIMILARITY.
FT DISULFID 328 340 BY SIMILARITY.
FT DISULFID 351 370 BY SIMILARITY.
FT CARBOHYD 82 82 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 127 127 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 169 169 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 205 205 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 294 294 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 529 529 N-LINKED (GLCNAC. .) (POTENTIAL).
FT VARSPPLIC 402 433 VSPSYKIVARVDEPQNGQPIHGHGMLHAKQ -> RLINS
FT FT CBYKRLKAGAPAPERQAEASTVTP (in isoform
FT FT LMW)
FT FT /FTid-VSP 001265.
FT FT Missing (in isoform LMW).
FT FT /FTid-VSP 001266.
FT FT E -> K (IN REF. 2).
FT SEQUENCE 639 AA; 70933 MW; D3172DF94FF56AF5 CRC64;
Query Match 59.7%; Score 410; DB 1; Length 639;
Best Local Similarity 66.4%; Pred. No. 2e-31; Indels 0; Gaps 0;
Matches 77; Conservative 13; Mismatches 26;
Qy 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFYFKIDNVKKEARVQV 62
Db 253 GDDLFELLPEDCPGCPNIPVDSPELKEALGHSIAQLNAENHHTFYFKIDTVKXATSQV 312
Qy 63 AKKYPIDFVARETCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYTV 118
Db 313 AGTKYVIEFIARETCSKESNAELTADCTKRLGQSLNCNANVYMRPWENKVPV 368
RESULT 8
KNT2_RAT
ID KNT2_RAT STANDARD; PRT; 430 AA.
AC P08932;
DT 01-NOV-1988 (Rel. 09, Created)
DT 01-NOV-1988 (Rel. 09, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE T-kininogen II precursor (Major acute phase protein) (Alpha-1-MAP)
DE (Thiostatin) [Contains: T-kinin].
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

```

```

OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=86008264; PubMed=2413018;
RA Furuto-Kato S., Matsumoto A., Kitamura N., Nakanishi S.;
RT "Primary structures of the mRNAs encoding the rat precursors for
RT bradykinin and T-kinin. Structural relationship of kininogens with
RT major acute phase protein and alpha 1-cysteine proteinase
RT inhibitor.";
RL J. Biol. Chem. 260:12054-12059 (1985).
CC -I- FUNCTION: Kininogens are plasma glycoproteins with a number of
CC functions: (1) as precursor of the active peptide bradykinin they
CC effect smooth muscle contraction, induction of hypotension and
CC increase of vascular permeability. (2) They play a role in blood
CC coagulation by helping to position optimally prekallikrein and
CC factor XI next to factor XII. (3) They are inhibitor of thiol
CC proteases.
CC -I- SUBCELLULAR LOCATION: Extracellular.
CC -I- TISSUE SPECIFICITY: Plasma.
CC -I- INDUCTION: In response to an inflammatory stimulant. T-kininogen
CC II synthesis is induced and the plasma concentration of
CC T-kininogen I is raised.
CC -I- PTM: AS T-KININ IS PRECEDED BY A MET INSTEAD OF AN ARG OR LYS, IT
CC IS NOT RELEASED FROM ITS PRECURSOR BY EITHER TISSUE OR PLASMA
CC KALLIKREIN.
CC -I- MISCELLANEOUS: Rats express four types of kininogens: the
CC classical HMW and LMW kininogens produced by alternative splicing
CC of the same gene, and two additional LMW-like kininogens: T-I and
CC T-II.
CC -I- SIMILARITY: Contains 3 cystatin-like domains.
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CC or send an email to license@isb-sib.ch).
CC EMBL; M11885; AAA41491.1; -
DR PIR; B28055; B28055.
DR GlycoSuiteDB; P08932; -.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 3.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;
KW Thiol protease inhibitor; Bradykinin; Acute phase; Signal.
FT SIGNAL 1 18
FT CHAIN 19 430 KININOGEN, T-II.
FT CHAIN 19 375 HEAVY CHAIN.
FT PEPTIDE 376 386 T-KININ.
FT CHAIN 387 430 LIGHT CHAIN.
FT DOMAIN 19 135 CYSTATIN-LIKE 1.
FT DOMAIN 136 257 CYSTATIN-LIKE 2.
FT DOMAIN 258 375 CYSTATIN-LIKE 3.
FT DISULFID 28 404 INTERCHAIN (BY SIMILARITY).
FT DISULFID 83 94 BY SIMILARITY.
FT DISULFID 107 125 BY SIMILARITY.
FT DISULFID 141 144 BY SIMILARITY.
FT DISULFID 205 217 BY SIMILARITY.
FT DISULFID 228 247 BY SIMILARITY.
FT DISULFID 263 266 BY SIMILARITY.
FT DISULFID 327 339 BY SIMILARITY.
FT DISULFID 350 369 BY SIMILARITY.
FT CARBOHYD 82 82 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 126 126 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 326 326 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 430 AA; 47524 MW; 43EDF02D1BF55076 CRC64;
Query Match 56.5%; Score 388; DB 1; Length 430;

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Best Local Similarity 62.1%; Pred. No. 1.6e-29;		Matches 72; Conservative 15; Mismatches 29; Indels 0; Gaps 0;	
Qy	3 GKQVQPTKICVCPDIPNSPELSEETLTHITKLNAENNAATFYFKIDNVKARQVV 62		
Dy	252 GDDUFSLLPKKFCPCPKNIPVDSPELKEALCHSIAQLNAGNHLFFKIDTVKATSOVV 311		
Qy	63 AGKGYFDVARETTCESKEEELTESCTKKLGQSLDCNAEVVVPWEKKIYPTV 118		
Dy	312 AGTKYVIEFTARETNCSTQNTLTADCTCKHLGQSLNCNANVYMPWENKVPTV 367		
RESULT 9			
KNTL1 RAT			
AC	P01048; P04081; STANDARD; PRT; 430 AA.		
DT	01-NOV-1986 (Rel. 03, Created)		
DT	01-NOV-1988 (Rel. 09, Last sequence update)		
DT	15-MAR-2004 (Rel. 43, Last annotation update)		
DE	T-kininogen I precursor (Major acute phase protein) (Alpha-1-MAP)		
DE	(Thioistatin) [Contains: T-kinin].		
GN	MAP1.		
OS	Rattus norvegicus (Rat).		
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.		
OK	NCBI_TaxID=10116;		
RP	SEQUENCE FROM N.A.		
RA	MEDLINE=86008264; PubMed=2413018;		
RA	Furuto-Kato S., Matsumoto A., Kitamura N., Nakanishi S.;		
RT	"Primary structures of the MRNAs encoding the rat precursors for		
RT	bradykinin and T-kinin. Structural relationship of kininogens with		
RT	major acute phase protein and alpha 1-cysteine proteinase		
RT	inhibitor."		
RL	J. Biol. Chem. 260:12054-12059(1985).		
RP	SEQUENCE OF 5-430 FROM N.A., AND PARTIAL SEQUENCE.		
RP	MEDLINE=86008266; PubMed=2413019;		
RA	Anderson K.P., Heath E.C.;		
RT	"The relationship between rat major acute phase protein and the		
RT	kininogens."		
RL	J. Biol. Chem. 260:12065-12071(1985).		
RP	SEQUENCE OF 7-430 FROM N.A.		
RP	MEDLINE=85127561; PubMed=2578992;		
RA	Cole T., Inglis A.S., Roxburgh C.M., Howlett G.J., Schreiber G.;		
RT	"Major acute phase alpha 1-protein of the rat is homologous to bovine		
RT	kininogen and contains the sequence for bradykinin: its synthesis is		
RT	regulated at the mRNA level."		
RL	FEBS Lett. 182:57-61(1985).		
RP	SEQUENCE OF 1-65 FROM N.A.		
RA	Fung W.-P., Schreiber G.;		
RT	"Structure and expression of the genes for major acute phase alpha 1-		
RT	protein (thioistatin) and kininogen in the rat."		
RL	J. Biol. Chem. 262:9298-9308(1987).		
CC	1- FUNCTION: Kininogens are plasma glycoproteins with a number of		
CC	functions: (1) as precursor of the active peptide bradykinin they		
CC	effect smooth muscle contraction, induction of hypotension and		
CC	increase of vascular permeability. (2) they play a role in blood		
CC	coagulation by helping to position optimally prekallikrein and		
CC	factor XI next to factor XII. (3) They are inhibitor of thiol		
CC	proteases.		
CC	1- SUBCELLULAR LOCATION: Extracellular.		
CC	1- TISSUE SPECIFICITY: Plasma		
CC	1- INDUCTION: In response to an inflammatory stimulant. T-kininogen		
CC	II synthesis is induced and the plasma concentration of		
CC	T-kininogen I is raised.		
CC	1- PTM: AS T-KININ IS PRECEDED BY A MET INSTEAD OF AN ARG OR LYS, IT		
CC	IS NOT RELEASED FROM ITS PRECURSOR BY EITHER TISSUE OR PLASMA		
CC	KALLIKREIN		
CC	1- MISCELLANEOUS: Rats express four types of kininogens: the		

CC	classical HMW and LMW kininogens produced by alternative splicing		
CC	of the same gene, and two additional LMW-like kininogens: T-I and		
CC	T-II.		
CC	1- SIMILARITY: Contains 3 cystatin-like domains.		
CC	1- CAUTION: In addition to the conflicts described in the feature		
CC	table, Ref.2 sequence differs from that shown in positions 257,		
CC	262, 268, 269, 295, 314, 315, 331, 332 and 389. In all those		
CC	positions the alternate amino acid is the one present in T-II		
CC	kininogen.		
CC	-----		
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CC	between the Swiss Institute of Bioinformatics and the EMBL outstation -		
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CC	use by non-profit institutions as long as its content is in no way		
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CC	entities requires a license agreement (See http://www.isb-sib.ch/announce/		
CC	or send an email to license@isb-sib.ch).		
CC	-----		
CC	EMBL; M11803; AAA41489.1; -		
DR	EMBL; M11661; AAA41570.1; -		
DR	EMBL; M16454; AAA41568.1; -		
DR	EMBL; X02239; CAA326182.1; ALT_SEQ.		
DR	PIR; A01286; KGRTH.		
DR	PIR; A23897; A23897.		
DR	PIR; A27115; A27115.		
DR	GlycoSuiteDB; P01048; -		
DR	InterPro; IPR000010; Cystatin.		
DR	Pfam; PF00031; Cystatin; 3.		
DR	SMART; SM00043; CY_3		
DR	PROSITE; PS00287; CYSTATIN; 2.		
KW	Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;		
KW	Thiol protease inhibitor; Bradykinin; Acute phase; Signal.		
FT	SIGNAL 1 18		
FT	CHAIN 19 430 KININOGEN, T-I.		
FT	CHAIN 19 375 HEAVY CHAIN.		
FT	PEPTIDE 376 386 T-KININ.		
FT	CHAIN 387 430 LIGHT CHAIN.		
FT	DOMAIN 19 135 CYSTATIN-LIKE 1.		
FT	DOMAIN 136 257 CYSTATIN-LIKE 2.		
FT	DOMAIN 258 375 CYSTATIN-LIKE 3.		
FT	DISULFID 28 404 INTERCHAIN (BY SIMILARITY).		
FT	DISULFID 83 94 BY SIMILARITY.		
FT	DISULFID 107 125 BY SIMILARITY.		
FT	DISULFID 141 144 BY SIMILARITY.		
FT	DISULFID 205 217 BY SIMILARITY.		
FT	DISULFID 228 247 BY SIMILARITY.		
FT	DISULFID 263 266 BY SIMILARITY.		
FT	DISULFID 327 339 BY SIMILARITY.		
FT	DISULFID 350 369 BY SIMILARITY.		
FT	CARBOHYD 82 82 N-LINKED (GLCNAC.) (POTENTIAL).		
FT	CARBOHYD 126 126 N-LINKED (GLCNAC.) (POTENTIAL).		
FT	CARBOHYD 168 168 N-LINKED (GLCNAC.) (POTENTIAL).		
FT	CARBOHYD 204 204 N-LINKED (GLCNAC.) (POTENTIAL).		
FT	CARBOHYD 326 326 N-LINKED (GLCNAC.) (POTENTIAL).		
FT	CONFLICT 26 28 LNC -> MDR (IN REF. 2).		
FT	CONFLICT 55 55 V -> L (IN REF. 2).		
FT	CONFLICT 61 61 E -> K (IN REF. 1).		
FT	CONFLICT 83 83 C -> Y (IN REF. 3).		
FT	CONFLICT 166 166 REV -> TKI (IN REF. 2).		
FT	CONFLICT 179 181 S -> F (IN REF. 2 AND 3).		
FT	CONFLICT 193 193 N -> D (IN REF. 2).		
FT	CONFLICT 212 212 S -> F (IN REF. 2).		
FT	CONFLICT 214 214 R -> H (IN REF. 3).		
FT	CONFLICT 229 229 T -> R (IN REF. 2).		
FT	CONFLICT 233 233 H -> Y (IN REF. 2).		
FT	CONFLICT 257 257 E -> S (IN REF. 2).		
FT	CONFLICT 262 262 N -> K (IN REF. 2).		
FT	CONFLICT 264 264 R -> F (IN REF. 2).		
FT	CONFLICT 268 268 RE -> KN (IN REF. 2).		
FT	CONFLICT 295 295 I -> L (IN REF. 2).		
FT	CONFLICT 314 315 VI -> TK (IN REF. 2).		
FT	CONFLICT 331 332 SK -> TN (IN REF. 2).		
FT	CONFLICT 389 389 R -> Q (IN REF. 2).		

RA Levaeslao M.H., Leversha M.A., Lloyd C., Lloyd D.M., Lovell J.D.,
RA Marsh V.L., Martin S.D., McConachie L.J., McIay K., McMurray A.A.,
RA Milne S.L., Misry D., Moore M.J.F., Mullikin J.C., Nickerson T.,
RA Oliver K., Parker A., Patel R., Pearce T., Plumb R.W., Ramsay H.,
RA Phillimore B.J.C.T., Prathalingam S.R., Plumb R.W., Shownkeen R., Sims S.,
RA Rice C.M., Ross M.T., Scott C.E., Saha H.K., Steward C.A., Sulston J.E.,
RA Skuce C.D., Smith M.L., Soderlund C., Steward C.A., Sulston J.E.,
RA Swann R.M., Sycamore A.C., Taylor R., Tee L., Thomas D.W., Thorpe A.,
RA Tracey A., Tromans A.C., Vaudin M., Wall M., Wallis J.M., Williams S.A.,
RA Whitehead S.L., Whittaker P., Willey D.L., Williams L., Williams S.A.,
RA Wilming L., Wray P.W., Hubbard T., Durbin R.M., Bentley D.R., Beck S.,
RA Rogers J.;
RT "The DNA sequence and comparative analysis of human chromosome 20.";
RL Nature 414:865-871(2001).
RN [6].
RP SEQUENCE FROM N.A.
RC TISSUE=Lung;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S.J., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Farley J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting J., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smallos D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -!- FUNCTION: Inhibits papain and cathepsin L but with affinities
CC lower than other cystatins. May play a role in immune regulation
CC through inhibition of a unique target in the hematopoietic system.
CC -!- SUBCELLULAR LOCATION: Secreted (Probable).
CC -!- TISSUE SPECIFICITY: Primarily expressed in peripheral blood cells
CC and spleen.
CC -!- SIMILARITY: Belongs to the cystatin family.
CC
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CC
CC EMBL; AF036342; AAC35747.1; -;
CC EMBL; AF031824; AAC39788.1; -;
CC EMBL; AB015225; BAA34941.1; ALT_INIT.
CC EMBL; AB029636; BAB11886.1; ALT_INIT.
CC EMBL; AL035661; CAB75498.1; -;
CC EMBL; BC015507; AAB15507.1; ALT_INIT.
CC HSSP; P01034; IG96.
CC Genew; HGNC:2479; CST7.
CC NIM; 603253; -;
CC GO; GO:0004869; F:cysteine protease inhibitor activity; TAS.
CC GO; GO:0006955; P:immune response; TAS.
CC InterPro; IPR000010; Cystatin.
CC Pfam; PFC00031; cystatin; 1.
CC SMART; SM00043; Cy; 1.
CC PROSITE; PS00287; CYSTATIN; 1.
CC Thiol protease inhibitor; Glycoprotein; Signal.
CC SIGNAL 1 19 POTENTIAL.
CC CHAIN 20 145 CYSTATIN F.
CC ACT_SITE 37 37 REACTIVE SITE.

FT SITE 81 85 SECONDARY AREA OF CONTACT.
FT DISULFID 99 110 BY SIMILARITY.
FT DISULFID 124 144 BY SIMILARITY.
FT CARBOHYD 62 62 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 115 115 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 145 AA; 16454 MW; B2BCC4F76857CB0F CRC64;
Query Match 23.8%; Score 163.5; DB 1; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.9e-09;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;
QY 11 TKICVCGREDIPTNSPELEETLTHITKLNNATVFYFKIDNVKKARVQVVGKKYFID 70
DB 32 SRVKFGPPTKINTNDPGVLQAAARYSVKFNCTNDFLFXESRITFALVQIVKGLKYLE 91
QY 71 FVARETTCKSKESNEBELTESCE---TKKLGQSLDCAEAVVYVFWKPKIYPTVTVNWE 124
DB 92 VEIGRTTCKKNQHLRL-DDCDFQTNHLTKQLTLCYSEVVVWPW-----LQHFE 138
RESULT 12
CYTC_MACMU STANDARD; PRT; 146 AA.
AC Q19092;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Cystatin C precursor.
GN CST3.
OS Macaca mulatta (Rhesus macaque).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopitheidae;
OC Cercopithecinas; Macaca.
OX NCBI_TaxID=9544;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97054523; PubMed=8988820;
RA Wei L.H., Walker L.C., Levy E.;
RT "Cystatin C. Icelandic-like mutation in an animal model of
RT cerebrovascular beta-amyloidosis.";
RL Stroke 27:2080-2085(1996).
CC -!- FUNCTION: As an inhibitor of cysteine proteinases, this protein is
CC thought to serve an important physiological role as a local
CC regulator of this enzyme activity.
CC -!- SIMILARITY: Belongs to the cystatin family.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; U51912; AAB64050.1; -;
CC HSSP; P01034; IG96.
CC InterPro; IPR000010; Cystatin.
CC Pfam; PFC00031; cystatin; 1.
CC SMART; SM00043; Cy; 1.
CC PROSITE; PS00287; CYSTATIN; 1.
CC Thiol protease inhibitor; Amyloid; Signal.
CC SIGNAL 1 26
CC CHAIN 27 146 CYSTATIN C.
CC ACT_SITE 37 37 REACTIVE SITE.
CC SITE 81 85 SECONDARY AREA OF CONTACT.
CC DISULFID 99 109 BY SIMILARITY.
CC DISULFID 123 143 BY SIMILARITY.
CC SEQUENCE 146 AA; 15857 MW; F0B3BB774A29DF26 CRC64;
Query Match 20.2%; Score 138.5; DB 1; Length 145;
Best Local Similarity 27.9%; Pred. No. 2.4e-06;
Matches 34; Conservative 25; Mismatches 52; Indels 11; Gaps 4;

QY 8 OPTKICVGPDPPTNSPELEBTLTHITIKLNAENNAFYKIDNVKKARVQVAGKY 67
 Db 31 KPRP-LVGPMDASVEEGREALDFAVSEYNKASNDYHSRALQVPRKQIVAGVNY 88
 QY 68 FIDFVARETTCSKESNEELTESC---ETKLGQSLDCNAEVVVPWEKKIYPVTVNHWE 124
 Db 89 FLDELGRITCTK--TQPNLDCNPFHEQPHLKKKACSPQIYTVPWQ-----GTWLSKST 142
 QY 125 CE 126
 Db 143 CQ 144

RESULT 13
 CYTM_HUMAN STANDARD; PRT; 149 AA.
 ID CYTM_HUMAN
 AC Q15828;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Cystatin M precursor (Cystatin E).
 GN CST6.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97150844; PubMed=8995380;
 RA Sotiropoulou G., Anisowicz A., Sager R.;
 RT "Identification, cloning, and characterization of cystatin M, a novel
 RT cysteine proteinase inhibitor, down-regulated in breast cancer.";
 RL J. Biol. Chem. 272:903-910(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97256812; PubMed=9099741;
 RA Ni J., Abrahamson M., Zhang M., Fernandez M.A., Grubb A., Su J.,
 RA Yu G.L., Li Y., Parmelee D., Xing L., Coleman T.A., Gentz S.,
 RA Thotakura R., Nguyen N., Hesselberg M., Gentz R.;
 RT "Cystatin E is a novel human cysteine proteinase inhibitor with
 RT structural resemblance to family 2 cystatins.";
 RL J. Biol. Chem. 272:10853-10858(1997).
 RN [3]
 RP SEQUENCE FROM N.A.
 RX TISSUE=Prostate;
 RC MEDLINE=2338257; PubMed=12477932;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Ustin T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahey J., Helton E., Kettner M., Madan A., Rodrigues S., Sanchez A.,
 RA Blakesley R.W., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Rodriguez A.C., Touchman J.W., Green E.D., Dickson M.C.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalusz D.B.,
 RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [4]
 RP CHARACTERIZATION, AND TISSUE SPECIFICITY.
 RX MEDLINE=21246880; PubMed=11348457;
 RA Zeeuwen P.L., Van Vlijmen-Willems I.M., Jansen B.J., Sotiropoulou G.,
 RA Curfs J.H., Meis J.F., Janssen J.J., Van Ruissen F., Schalkwijk J.;
 RT "Cystatin M/E expression is restricted to differentiated epidermal
 RT keratinocytes and sweat glands: a new skin-specific proteinase

RT inhibitor that is a target for cross-linking by transglutaminase.";
 RL J. Invest. Dermatol. 116:693-701(2001).
 CC -!- FUNCTION: Shows moderate inhibition of cathepsin B but is not
 CC active against cathepsin C.
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- TISSUE SPECIFICITY: Restricted to the stratum granulosum of normal
 CC skin, the stratum granulosum/spinosum of psoriatic skin, and the
 CC secretory coils of eccrine sweat glands. Low expression levels are
 CC found in the nasal cavity.
 CC -!- PTM: Substrate for transglutaminases. Acts as an acyl acceptor but
 CC not as an acyl donor.
 CC -!- SIMILARITY: Belongs to the cystatin family.
 CC
 CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
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 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC
 CC -----
 CC EMBL; U62800; RAB06566.1; -;
 CC EMBL; U81233; RAB61305.1; -;
 CC EMBL; BC011334; AAH31334.1; -;
 CC HSP; P01038; ICEW
 CC Genew; HGNC:2478; CST6.
 CC MIM; 601891; -;
 CC GO; GO:0004869; F:cysteine protease inhibitor activity; TAS.
 CC GO; GO:0007345; P:embryogenesis and morphogenesis; TAS.
 CC InterPro; IPR000010; Cystatin.
 CC Pfam; PF00031; cystatin; 1.
 CC SMART; SM00043; CY; 1
 CC PROSITE; PS00287; CYSTATIN; 1.
 CC Thiol protease inhibitor; Signal; Glycoprotein.
 KW SIGNAL 1 28 PROBABLE.
 FT CHAIN 29 149 CYSTATIN M.
 FT ACT SITE 36 36 REACTIVE SITE.
 FT SITE 80 84 SECONDARY AREA OF CONTACT.
 FT DISULFID 98 113 BY SIMILARITY.
 FT DISULFID 126 146 BY SIMILARITY.
 FT CARBOHYD 137 137 N-LINKED (GLCNAC...) (POTENTIAL).
 SQ SEQUENCE 149 AA; 16511 MW; 2076A78BFC9FAC8C CRC64;
 Query Match 20.2%; Score 138.5; DB 1; Length 149;
 Best Local Similarity 31.5%; Pred. No. 2.4e-06;
 Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
 QY 10 PTKICVGPDPPTNSPELEBTLTHITIKLNAENNAFYKIDNVKKARVQVAGKYPT 69
 Db 30 PQERMYGELRLSPDPQVQKAAQAAVASYNMGNSIYYFRDTHIKAQSLVAGIKYFL 89
 QY 70 DFVARETTCSKE---SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
 Db 90 TWEMGSTDCRKTREVTGDHVDLT--TCLAAGAQQEKLRCDFEVLVVPWQ 136
 RESULT 14
 ID CYTC_BOVIN STANDARD; PRT; 148 AA.
 CYTC_BOVIN
 AC P01035;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 15-JUL-1999 (Rel. 38, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Cystatin C precursor (Colostrum thiol proteinase inhibitor).
 GN CST3.
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.; SEQUENCE OF 66-83, AND CHARACTERIZATION.
 RC TISSUE=Cerebrospinal fluid, and Choroid plexus;

RA MEDLINE=98094199; PubMed=9434110;
RA Olsson S.-L., Ek B., Wilm M., Broberg S., Rask L., Bjoerk I.;
RA "Molecular cloning and N-terminal analysis of bovine cystatin C
RA identification of a full-length N-terminal region.";
RL Biochim. Biophys. Acta 1343:203-210(1997).
RN (2)
RP SEQUENCE OF 37-148.
RX MEDLINE=85231205; PubMed=3891407;
RX Hirado M., Tsunawasa S., Sakiyama F., Niinobe M., Fujii S.;
RA "Complete amino acid sequence of bovine colostrum low-Mr cysteine
RA proteinase inhibitor.";
RL FEBS Lett. 186:41-45(1985).
CC -!- FUNCTION: This is a thiol proteinase inhibitor.
CC -!- MASS SPECTROMETRY: MW=13420; METHOD=MALDI.
CC -!- SIMILARITY: Belongs to the cystatin family.
CC
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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC use by non-profit institutions as long as its content is in no way
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CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; Y10811; CAA71771.1; -.
DR HSP; P01034; 1G96.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00287; CYSTATIN; 1.
KW Thiol protease inhibitor; Signal: Pyrrolidone carboxylic acid.
FT SIGNAL 1 30
FT CHAIN 31 148 CYSTATIN C.
FT MOD RES 31 31 PYRROLIDONE CARBOXYLIC ACID (PROBABLE).
FT ACT SITE 40 40 REACTIVE SITE.
FT SITE 84 88 SECONDARY AREA OF CONTACT.
FT DISULFID 102 112 BY SIMILARITY.
FT DISULFID 126 146 BY SIMILARITY.
SQ SEQUENCE 148 AA; 16265 MW; EE740FE37CFB9FOE CRC64;
Query Match 20.0%; Score 137.5; DB 1; Length 148;
Best Local Similarity 28.8%; Pred. No. 3e-06;
Matches 32; Conservative 25; Mismatches 35; Indels 19; Gaps 4;
OY 24 NSPELEETLTIYIKLNENATYFFKIDNVKARQVWAGKYFIDFVARETTCSKESN 83
DB 48 NEEGVQEALSPAVSEFNKRSDAYQSRVYVRARQVWGMVYFLDVLGRITCTK--S 105
OY 84 EELTESC-----ETKKLGSLDCNAEVVVPWEKKIYPTVTNHWCE 126
DB 106 QANLDCPFHQPHLKREKL-----CSFQVYVPMN-----TINLVKFSQ 147
RESULT 15
ID FETB RAT STANDARD; PRT; 378 AA.
AC Q9OX79.
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Fetusin-B precursor (IRL685).
GN FETUB.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Sprague-Dawley; TISSUE=Liver;
RX MEDLINE=20407138; PubMed=10947975;
RA Olivier E., Soury E., Ruminy P., Husson A., Parmentier P., Daveau M.,
RA Sallier J.-P.;
RA "Fetusin-B, a second member of the fetusin family in mammals.";

RL Biochem. J. 350:589-597(2000).
CC -!- SUBCELLULAR LOCATION: Secreted (Potential).
CC -!- TISSUE SPECIFICITY: Liver.
CC -!- SIMILARITY: Belongs to the fetusin family.
CC -!- SIMILARITY: Contains 2 cystatin-like domains.
CC
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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AJ24926; CAB62543.1; -.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; CY; 2.
DR PROSITE; PS01254; FETUIN_1; 1.
DR PROSITE; PS01255; FETUIN_2; 1.
KW Glycoprotein; Signal; Repeat.
FT SIGNAL 1 18 POTENTIAL.
FT CHAIN 19 378 FETUIN-B.
FT DOMAIN 27 152 CYSTATIN-LIKE 1.
FT DOMAIN 153 273 CYSTATIN-LIKE 2.
FT DISULFID 96 107 BY SIMILARITY.
FT DISULFID 120 140 BY SIMILARITY.
FT DISULFID 154 157 BY SIMILARITY.
FT DISULFID 217 224 BY SIMILARITY.
FT DISULFID 237 260 BY SIMILARITY.
FT CARBOHYD 40 40 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 139 139 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 378 AA; 41532 MW; 066C0A5C3B03C878 CRC64;
Query Match 19.7%; Score 135; DB 1; Length 378;
Best Local Similarity 25.8%; Pred. No. 1.5e-05;
Matches 33; Conservative 32; Mismatches 49; Indels 14; Gaps 5;
OY 7 VQPTK-----ICVGPDRIPNTPNSPELETLTHITKLNENATYFFKIDNVKARQV 61
DB 142 LRPVSKRKTHSMCPDCPHFVDLSAPSVLEAATESLAKFNSENPCKQYALV-KVTKATTOW 200
OY 62 VAGKTYFIDFVARETTCSKESNEELTESCTETKKLGSLDCNAEVVVPW-EKKIYPTVT 119
DB 201 VVGPSYFVEYLKESPTQSDSCSLQASDSEPVGL---CQGLSKSPGVPPQRFKKTWT 257
OY 120 VNHWECEP 127
DB 258 VS---CEF 262

Search completed: September 24, 2004, 14:09:13

Job time : 9.636 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:06:08 ; Search time 13.716 Seconds
(without alignments)
890.662 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687

Sequence: 1 GSGKDFVQPTKICVGCPRD.....VPWEKKIYPTVTWNHWECEP 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR.78.*

1: Pir1.*
2: Pir2.*
3: Pir3.*
4: Pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	618	90.0	427	1 KGHUL1	kininogen, LMW pre
2	618	90.0	444	1 KGHUL1	kininogen, LMW pre
3	440	64.0	436	1 KGBOL1	kininogen, LMW I p
4	440	64.0	621	1 KGBOL1	kininogen, LMW I p
5	413	60.1	434	1 KGBOL2	kininogen, LMW II
6	413	60.1	619	1 KGBOL2	kininogen, LMW II
7	410	59.7	433	2 A28055	K-kininogen, LMW I
8	410	59.7	639	2 A25486	kininogen, LMW I p
9	388	56.5	430	2 A23897	major acute phase
10	388	56.5	430	2 B28055	T-kininogen, LMW I
11	381	55.5	423	1 KGRTM	major acute phase
12	380	55.3	430	1 KGRTM	T-kininogen I prec
13	137.5	20.0	146	1 UDBO	cystatin - bovine
14	132.5	19.3	146	1 UDBO	cystatin C precurs
15	130	18.9	127	2 S07085	cystatin C precurs
16	129	18.8	120	2 S10587	cystatin C - rat
17	128	18.6	111	2 A28793	cystatin - puff ad
18	127.5	18.6	140	2 A36163	cystatin C precurs
19	124.5	18.1	141	2 B29632	cystatin SA precur
20	118.5	17.2	139	1 UDCH	cystatin precursor
21	113	16.4	141	2 JQ1470	cystatin S precurs
22	112	16.3	91	2 S68034	T-kininogen (clone
23	112	16.3	91	2 S68035	T-kininogen (clone
24	111	16.2	111	1 JC2040	cystatin - chum sa
25	109.5	15.9	141	1 UDHP2	cystatin SN precur
26	108.5	15.8	141	1 UDHP1	cystatin S precurs
27	107	15.6	139	2 T33740	hypothetical prote
28	106	15.4	132	2 JC4918	cystatin precursor
29	105.5	15.4	162	2 A43428	onchocystatin - ne

RESULT 1

KGHUL1

kininogen, LMW precursor [validated] - human

N;Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen

N;Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen

C;Species: Homo sapiens (man)

C;Date: 06-Jul-1982 #sequence revision 27-Nov-1985 #text change 08-Dec-2000

C;Accession: A01280; B25276; A27900; A27699; A31905; A34030

R;Ohkubo, I.; Kurachi, K.; Takasawa, T.; Shiohara, H.; Sasaki, M.

Biochemistry 23, 5691-5697, 1984

A;Title: Isolation of a human cDNA for alpha-2-thiol proteinase inhibitor and its identi-

A;Reference number: A90490; MUID:85122621; PMID:6441591

A;Accession: A01280

A;Molecule type: mRNA

A;Residues: 1-427 <CHK>

A;Cross-references: GB:K02566; NID:g177889; PIDN:AAA35497.1; PID:g177890

R;Takagaki, Y.; Kitamura, N.; Nakanishi, S.

J. Biol. Chem. 260, 8601-8609, 1985

A;Title: Cloning and sequence analysis of cDNAs for human high molecular weight and low

A;Reference number: A92544; MUID:85234582; PMID:2989293

A;Accession: B25276

A;Molecule type: mRNA

A;Residues: 1-427 <TAK>

A;Cross-references: GB:M11437; NID:g186751; PIDN:AAB59551.1; PID:g386853

R;Lottspeich, F.; Kellermann, J.; Henschen, A.; Rauth, G.; Mueller-Esterl, W.

in Kinins IV, part A, Greenbaum, L.M., and Margolius, H.S., eds., pp.91-95, Plenum, New

A;Title: Amino acid sequence of the light chain of human low molecular mass kininogen.

A;Reference number: A27900

A;Accession: A27900

A;Molecule type: protein

A;Residues: 390-427 <LOT>

R;Mindrou, T.; Carretero, O.A.; Proud, D.; Walz, D.; Scicli, A.G.

Biochem. Biophys. Res. Commun. 152, 519-526, 1988

A;Title: A new kinin moiety in human plasma kininogens.

A;Reference number: A27699; MUID:88209021; PMID:3365237

A;Accession: A27699

A;Molecule type: protein

A;Residues: 380-389 <MIN>

R;Maeda, H.; Matsumura, Y.; Kato, H.

J. Biol. Chem. 263, 16051-16054, 1988

A;Title: Purification and identification of [hydroxyprolyl(3)]-bradykinin in ascitic fluid

A;Reference number: A31905; MUID:89034061; PMID:3182782

A;Accession: A31905

A;Molecule type: protein

A;Residues: 381-389 <MAE>

R;Sasaguri, M.; Ikeda, M.; Ideishi, M.; Arakawa, K.

Biochem. Biophys. Res. Commun. 150, 511-516, 1988

A;Title: Identification of [hydroxyproline(3)]-lysyl-bradykinin released from human plas-

A;Reference number: A34030; MUID:88106632; PMID:3337729

A;Accession: A34030

A;Molecule type: protein

A;Residues: 380-389 <SAS>

A;Accession: A34030

A;Molecule type: protein

A;Residues: 380-389 <SAS>

A;Accession: A34030

A;Molecule type: protein

A;Residues: 380-389 <SAS>

R; Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
 J. Biol. Chem. 260, 8610-8617, 1985
 A; Title: Structural organization of the human kininogen gene and a model for its evolution
 A; Reference number: A92545; MUID: 85234583; PMID: 2989294
 A; Contents: annotation; gene organization
 R; Pierce, J. V., 52-57, 1968
 Fed. Proc. 27, 52-57, 1968
 A; Title: Structural features of plasma kinins and kininogens.
 A; Reference number: A91455; MUID: 90255622; PMID: 4952632
 A; Contents: annotation; bradykinin
 C; Comment: The LMW kininogen precursor is produced from the same gene as the HMW form (see A91455).
 C; Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
 C; Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, 1
 C; Comment: Bradykinin, released from kininogen prior to the release of bradykinin.
 C; Comment: A proline residue is present in the kininogen prior to the release of bradykinin.
 C; Genetics:
 A; Gene: GDB:KNG
 A; Cross-references: GDB:125256; OMIM:228960
 A; Map position: 3q27-3q27
 A; Introns: 65/3; 102/3; 131/1; 188/3; 224/3; 253/1; 310/3; 346/3; 375/3; 401/3
 C; Superfamily: kininogen; cystatin homology
 C; Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glyco
 F; 1-18/Domain: signal sequence #status predicted <SIG>
 F; 19-427/Product: LMW prokininogen (kininogen I) #status predicted <MAT>
 F; 19-389,390-427/Product: LMW kininogen II #status predicted <MAT2>
 F; 19-379/Product: LMW kininogen heavy chain #status predicted <HCH>
 F; 19-131/Domain: cystatin homology <CY1>
 F; 142-253/Domain: cystatin homology <CY2>
 F; 264-375/Domain: cystatin homology <CY3>
 F; 380-389/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>
 F; 381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
 F; 390-427/Product: LMW kininogen light chain #status experimental <LCH>
 F; 19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status predicted
 F; 28-407,83-94,107-126,142-145,206-218,229-248,264-267,328-340,351-370/Disulfide bonds:
 F; 48,469,205,294/Binding site: carboxylate (Asn) (covalent) #status predicted
 F; 379-380/Cleavage site: Met-Lys (kallikrein) #status experimental
 F; 383/Modified site: 4-hydroxyproline (Pro) (partial) #status experimental
 F; 389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
 F; 401/Binding site: carboxylate (Thr) (covalent) #status absent

Query Match 90.0%; Score 618; DB 1; Length 427;
 Best Local Similarity 100.0%; Pred. No. 3.6e-50;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTITIKLAENNAATFYFKIDNVKARQVQV 62
 DB 253 GKDFVQPTKICVGCPRDIPNTSPLEETLTITIKLAENNAATFYFKIDNVKARQVQV 312
 QY 63 AGKKYFIDFVARETTCSKESNEBELTSCETKKGQSLDCNDAEVYVPWEKKIYPTV 118
 DB 313 AGKKYFIDFVARETTCSKESNEBELTSCETKKGQSLDCNDAEVYVPWEKKIYPTV 368

RESULT 2
 KGHU1
 kininogen, HMW precursor [validated] - human
 N; Alternate names: alpha-2-thiol proteinase inhibitor; prokininogen; prokininogen
 N; Contains: bradykinin (kallidin I); HMW kininogen I; HMW kininogen II; low molecular we
 C; Species: Homo sapiens (man)
 C; Date: 28-May-1986 #sequence revision 28-May-1986 #text change 08-Dec-2000
 C; Accession: A01279; A25276; S32422; A91153; A24871; A27899; A27699; A31905; A34030; S02
 R; Ohkubo, I.; Kurachi, K.; Takasawa, T.; Shiohara, H.; Sasaki, M.
 Biochemistry 23, 5651-5697, 1984
 A; Title: Isolation of a human cDNA for alpha-2-thiol proteinase inhibitor and its ident
 A; Reference number: A90490; MUID: 85122621; PMID: 6441591
 A; Accession: A01279
 A; Molecule type: mRNA
 A; Residues: 1-389 <OHK>
 A; Cross-references: GB:K02566; MUID: g17789
 R; Takagaki, Y.; Kitamura, N.; Nakanishi, S.
 J. Biol. Chem. 260, 8601-8609, 1985
 A; Title: Cloning and sequence analysis of cDNAs for human high molecular weight and low
 A; Reference number: A92544; MUID: 85234582; PMID: 2989293
 A; Accession: A25276

A; Molecule type: mRNA
 A; Residues: 1-592, 'I', 594-644 <TAK>
 A; Cross-references: GB:M11437; MUID: g186751; PIDN: AAB59550.1; PID: g386852
 R; Auerwald, E. A.; Roessler, D.; Mentele, R.; Assfalg-Wachleideit, I.
 FEBS Lett. 321, 93-97, 1993
 A; Title: Cloning, expression and characterization of human kininogen domain 3.
 A; Reference number: S32422; MUID: 93223854; PMID: 8467916
 A; Accession: S32422
 A; Molecule type: mRNA
 A; Residues: 'ANSM', 253-377 <AUB>
 A; Note: differences are due to known cloning artifacts
 R; Lottspeich, F.; Kelleman, J.; Henschel, A.; Foerisch, B.; Muller-Esterl, W.
 Eur. J. Biochem. 152, 307-314, 1985
 A; Title: The amino acid sequence of the light chain of human high-molecular-mass kininogen
 A; Reference number: A91153; MUID: 86030270; PMID: 4054110
 A; Accession: A91153
 A; Molecule type: protein
 A; Residues: 379-644 <LOT>
 A; Note: the bradykinin sequence preceding the light chain sequence was not determined in
 R; Kelleman, J.; Lottspeich, F.; Henschel, A.; Mueller-Esterl, W.
 Eur. J. Biochem. 154, 471-478, 1986
 A; Title: Completion of the primary structure of human high-molecular-mass kininogen. The
 A; Reference number: A24871; MUID: 86108361; PMID: 3484703
 A; Accession: A24871
 A; Molecule type: protein
 A; Residues: 'Z', 20-380 <KEU>
 R; Kelleman, J.; Lottspeich, F.; Henschel, A.; Mueller-Esterl, W.
 in Kinins IV, Greenbaum, L.M., and Margolius, H.S., ed., pp. 85-89, Plenum Press, New York
 A; Title: Amino acid sequence of the light chain of human high molecular mass kininogen.
 A; Reference number: A27899
 A; Accession: A27899
 A; Molecule type: protein
 A; Residues: 379-389, K', 390-407, 'Q', 409-644 <KEL2>
 R; Mindrou, T.; Carrettero, O. A.; Proud, D.; Walz, D.; Scicli, A. G.
 Biochem. Biophys. Res. Commun. 152, 519-526, 1988
 A; Title: A new kinin moiety in human plasma kininogens.
 A; Reference number: A27699; MUID: 88209021; PMID: 3365237
 A; Accession: A27699
 A; Molecule type: protein
 A; Residues: 380-389 <MIN>
 R; Maeda, H.; Matsumura, Y.; Kato, H.
 J. Biol. Chem. 263, 16051-16054, 1988
 A; Title: Purification and identification of [hydroxyprolyl(3)]bradykinin in ascitic fluid
 A; Reference number: A31905; MUID: 89034061; PMID: 3182782
 A; Accession: A31905
 A; Molecule type: protein
 A; Residues: 381-389 <MAB>
 R; Sasaguri, M.; Ikeda, M.; Ideishi, M.; Arakawa, K.
 Biochem. Biophys. Res. Commun. 150, 511-516, 1988
 A; Title: Identification of [hydroxyproline(3)]-lysyl-bradykinin released from human plas
 A; Reference number: A34030; MUID: 88106632; PMID: 3337729
 A; Accession: A34030
 A; Molecule type: protein
 A; Residues: 380-389 <SAS>
 R; Lenarcic, B.; Gabrijelcic, D.; Rozman, B.; Drobnic-Kosorok, M.; Turk, V.
 Biol. Chem. Hoppe-Seyler 369, 257-261, 1988
 A; Title: Human cathepsin B and cysteine proteinase inhibitors (CPIs) in inflammatory and
 A; Reference number: S02482; MUID: 89076517; PMID: 3264507
 A; Accession: S02482
 A; Molecule type: protein
 A; Residues: 1-19,189-192; 310-314; 381-389 <LENI>
 R; Kato, H.; Matsumura, Y.; Maeda, H.
 FEBS Lett. 232, 252-254, 1988
 A; Title: Isolation and identification of hydroxyproline analogues of bradykinin in human
 A; Reference number: A61495; MUID: 88211869; PMID: 3366244
 A; Accession: A61495
 A; Molecule type: protein
 A; Residues: 380-389 <KAT1>
 A; Experimental source: urine
 A; Note: this peptide had Pro-383 modified to 4-hydroxyproline
 A; Accession: B61495
 A; Molecule type: protein
 A; Residues: 381-389 <KAT2>

A;Experimental source: urine
A;Note: this peptide had Pro-383 modified to 4-hydroxyproline
A;Accession: C61495
A;Molecule type: protein
A;Residues: 380-389 <KAT3>
R;Lenarcic, B.; Krasovec, M.; Ritonja, A.; Olafsson, I.; Turk, V.
FEBS Lett. 280, 211-215, 1991
A;Title: Inactivation of human cystatin C and kininogen by human cathepsin D.
A;Reference number: S14303; MUID:91192133; PMID:2013314
A;Accession: S14447
A;Molecule type: protein
A;Residues: 264-359, N'.361-375 <LEN2>
R;Little, S.S.; Johnson, D.A.
Biochem. J. 307, 341-346, 1995
A;Title: Human mast cell tryptase isoforms: separation and examination of substrate-specificity
A;Reference number: S55239; MUID:95251593; PMID:7733867
A;Accession: S55239
A;Molecule type: protein
A;Residues: 450-452, X', 454, X', 456 <LIT>
R;Straczek, J.; Naachi, F.; le Nguyen, D.; Becchi, M.; Heulin, M.H.; Nabet, P.; Bellevil
FEBS Lett. 373, 207-211, 1995
A;Title: Purification from human plasma of a tetrapeptide that potentiates insulin-like
A;Reference number: S68059; MUID:96033974; PMID:7599467
A;Accession: S68059
A;Molecule type: protein
A;Residues: 431-434 <STR>
R;Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 260, 8610-8617, 1985
A;Title: Structural organization of the human kininogen gene and a model for its evolution
A;Reference number: A92545; MUID:85234583; PMID:2989294
A;Contents: annotation; gene organization
R;Pierce, J.V.
Fed. Proc. 27, 52-57, 1968
A;Title: Structural features of plasma kinins and kininogens.
A;Reference number: A91455; MUID:90255622; PMID:4952632
A;Contents: annotation; bradykinin
C;Comment: The HMW kininogen precursor and the LMW form are produced from the same gene
C;Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C;Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is im
C;Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i
xyproline residue is present in the kininogen prior to the release of bradykinin.
C;Genetics:
A;Gene: GDB:KNG
A;Cross-references: GDB:125256; OMIM:228960
A;Map position: 3q27.3q27
A;Intons: 65/3; 102/3; 131/1; 188/3; 224/3; 253/1; 310/3; 346/3; 375/3
C;Superfamily: kininogen; cystatin homology
C;Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupl
F;19-644/Product: HMW kininogen I (prokininogen) #status experimental <MAT1>
F;19-379,390-644/Product: HMW kininogen II #status experimental <MAT2>
F;19-379/Domain: HMW kininogen heavy chain #status experimental <HCH>
F;19-379/Domain: HMW kininogen homology <CY1>
F;19-131/Domain: cystatin homology <CV2>
F;142-253/Domain: cystatin homology <CV3>
F;264-375/Domain: cystatin homology <CV3>
F;380-389/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>
F;381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
F;390-644/Domain: HMW kininogen light chain #status experimental <LCH>
F;421-510/Region: glycine/histidine/lysine-rich 30-residue repeats
F;431-434/Product: low molecular weight growth promoting factor #status experimental <GF
F;19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status experimen
F;28-614,83-94,107-126,142-145,206-218,229-248,264-267,328-340,351-370/Disulfide bonds:
F;48/Binding site: carboxylate (Asn) (covalent) #status absent
F;169,205,294/Binding site: carboxylate (Asn) (covalent) #status experimental
F;379-380/Cleavage site: Met-Lys (kallikrein) #status experimental
F;383/Modified site: 4-hydroxyproline (Pro) (partial) #status experimental
F;389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
F;401,533,542,546,557,571,593,628/Binding site: carboxylate (Thr) (covalent) #status ex
F;507/Binding site: carboxylate (Ser) (covalent) #status experimental
Query Match 90.0%; Score 618; DB 1; Length 644;
Best Local Similarity 100.0%; Pred. No. 5.7e-50; Indels 0; Gaps 0;
Matches 116; Conservative 0; Mismatches 0;

QY 3 GKDFVQPTKICVGCPRDIPNPSPELEETLTHITKLNAENNATFFKIDNVKKARQVW 62
DB 253 GKDFVQPTKICVGCPRDIPNPSPELEETLTHITKLNAENNATFFKIDNVKKARQVW 312
QY 63 AGKYFIDFVARETTCSKESNEELTESCTKLGQSLDCNAEVVVPWEKKIYPTV 118
DB 313 AGKYFIDFVARETTCSKESNEELTESCTKLGQSLDCNAEVVVPWEKKIYPTV 368
RESULT 3
KGBOL1
kininogen, LMW I precursor - bovine
N;Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N;Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C;Species: Bos primigenius taurus (cattle)
C;Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999
C;Accession: A01283
R;Nawa, H.; Kitamura, N.; Hirose, T.; Asai, M.; Inayama, S.; Nakanishi, S.
Proc. Natl. Acad. Sci. U.S.A. 80, 90-94, 1983
A;Title: Primary structures of bovine liver low molecular weight kininogen precursors and
A;Reference number: A91984; MUID:83117859; PMID:6572010
A;Accession: A01283
A;Molecule type: mRNA
A;Residues: 1-436 <NAW>
A;Cross-references: GB:J00010; GB:V00426; NID:G163256; PIDN:AAA30604.1; PID:G163257
C;Comment: The LMW kininogen precursor is produced from the same gene as the HMW form as
C;Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C;Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i
xyproline residue is present in the kininogen prior to the release of bradykinin.
C;Superfamily: kininogen; cystatin homology
C;Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glyci
F;1-18/Domain: signal sequence #status predicted <SIG>
F;19-436/Product: LMW kininogen I #status predicted <MAT>
F;19-378/Product: LMW kininogen I heavy chain #status predicted <HCH>
F;19-130/Domain: cystatin homology <CY1>
F;141-252/Domain: cystatin homology <CV2>
F;263-374/Domain: cystatin homology <CV3>
F;379-388/Product: lysyl-bradykinin (kallidin II) #status predicted <KBDY>
F;380-389/Product: bradykinin (kallidin I) #status predicted <BDY>
F;389-436/Product: LMW kininogen I light chain #status experimental <LCH>
F;19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status predicted
F;27-406,82-93,106-125,141-144,205-217,228-247,263-286,327-339,350-366/Disulfide bonds:
F;47,87,168,169,197,204/Binding site: carboxylate (Asn) (covalent) #status predicted
F;378-379/Cleavage site: Met-Lys (kallikrein) #status predicted
F;382/Modified site: 4-hydroxyproline (Pro) #status predicted
F;388-389/Cleavage site: Arg-Ser (kallikrein) #status predicted
Query Match 64.0%; Score 440; DB 1; Length 436;
Best Local Similarity 70.4%; Pred. No. 1.7e-33;
Matches 81; Conservative 14; Mismatches 20; Indels 0; Gaps 0;
QY 4 KDFVQPTKICVGCPRDIPNPSPELEETLTHITKLNAENNATFFKIDNVKKARQVWA 63
DB 253 KDFVQPTKICVGCPRDIPNPSPELEETLTHITKLNAENNATFFKIDNVKKARQVWA 312
QY 64 GKRYFIDFVARETTCSKESNEELTESCTKLGQSLDCNAEVVVPWEKKIYPTV 118
DB 313 GLKYSIVFTARETTCSKESNEELTESCTKLGQSLDCNAEVVVPWEKKIYPTV 367
RESULT 4
KGBOH1
kininogen, HMW I precursor - bovine
N;Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N;Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C;Species: Bos primigenius taurus (cattle)
C;Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999
C;Accession: A01281; A91923; A91938; A28559
R;Kitamura, N.; Takagaki, Y.; Furuto, S.; Tanaka, T.; Nawa, H.; Nakanishi, S.
Nature 305, 545-549, 1983
A;Title: A single gene for bovine high molecular weight and low molecular weight kininoge
A;Reference number: A93317; MUID:84014106; PMID:6571699

J. Biochem. 67, 313-323, 1970
A:Title: Studies on the structure of bovine kininogen: cleavages of disulfide bonds and
A:Reference number: A91923; MUID:70180420; PMID:4986212
A:Accession: A91923
A:Molecule type: protein
A:Residues: 376-391 <XAT>
F:Han, Y.N.; Kato, H.; Iwanaga, S.; Suzuki, T.
J. Biochem. 79, 1201-1222, 1976
A:Title: Primary structure of bovine plasma high-molecular-weight kininogen. The amino a
A:Reference number: A91941; MUID:76260155; PMID:956151
A:Accession: A91941
A:Molecule type: protein
A:Residues: 387-455 <HAN>
A:Note: 398-Pro, 401-Val, and 455-Lys were also found
F:Han, Y.N.; Kato, H.; Iwanaga, S.; Suzuki, T.
J. Biochem. 77, 55-68, 1975
A:Title: Studies on the primary structure of bovine high-molecular-weight kininogen. Ami
A:Reference number: A91938; MUID:75170265; PMID:1169237
A:Accession: A91938
A:Molecule type: protein
A:Residues: 456-496 <HA2>
F:Sueyoshi, T.; Miyata, T.; Hashimoto, N.; Kato, H.; Hayashida, H.; Miyata, T.; Iwanaga,
J. Biol. Chem. 262, 2768-2779, 1987
A:Title: Bovine high molecular weight kininogen. The amino acid sequence, positions of c
A:Reference number: A92627; MUID:87137530; PMID:3546295
A:Accession: B29559
A:Molecule type: protein
A:Residues: 'Z', 20-104, 'B', 106-256, 'XX', 257-376 <SUB>
R:Lottspeich, F.; Kellermann, J.; Henschen, A.; Foeretsch, B.; Muller-Esterl, W.
Eur. J. Biochem. 152, 307-314, 1985
A:Title: The amino acid sequence of the light chain of human high-molecular-mass kininog
A:Reference number: A91153; MUID:86030270; PMID:4054110
A:Contents: annotation; Bovine cleavage sites; bovine carbohydrate binding sites
F:Sueyoshi, T.; Miyata, T.; Kato, H.; Iwanaga, S.
Seikagaku 56, 808, 1984
A:Title: Disulfide bonds in bovine HMW kininogens.
A:Reference number: A94300
A:Contents: annotation; disulfide bonds
A:Note: article in Japanese
C:Comment: The HMW kininogen precursor is produced from the same gene as the LMW form as
C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C:Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is impo
C:Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i
xyproline residue is present in the kininogen prior to the release of bradykinin.
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupl
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-376/Product: HMW kininogen II #status predicted <MAT>
F:19-376/Product: HMW kininogen II heavy chain #status experimental <HCH>
F:19-130/Domain: cystatin homology <CY1>
F:141-253/Domain: cystatin homology <CY2>
F:261-372/Domain: cystatin homology <CY3>
F:377-386/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>
F:378-386/Product: bradykinin (kallidin I) #status experimental <BDY>
F:387-619/Product: HMW kininogen II light chain #status experimental <LCH>
F:418-488/Region: glycine/histidine/lysine-rich
F:19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status experimen
F:27-589, 82-93, 106-125, 141-144, 205-217, 228-247, 261-264, 325-337, 348-367/Disulfide bonds:
F:47/Binding site: carbohydrate (Asn) (covalent) #status absent
F:87, 168, 169, 204, 280/Binding site: carbohydrate (Asn) (covalent) #status experimental
F:136/Binding site: carbohydrate (Thr) (covalent) (partial) #status experimental
F:197/Binding site: carbohydrate (Asn) (covalent) (partial) #status experimental
F:376-377/Cleavage site: Met-Lys (kallikrein) #status experimental
F:380/Modified site: 4-hydroxyproline (Pro) #status predicted
F:386-387/Cleavage site: Arg-Ser (kallikrein) #status experimental
F:396, 400, 404, 510/Binding site: carbohydrate (Ser) (covalent) #status experimental
F:397-498, 516, 522, 534, 546, 555, 569/Binding site: carbohydrate (Thr) (covalent) #status ex
F:496-497/Cleavage site: Arg-Thr (kallikrein) #status experimental

QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNATFYFKIDNVKKARVQV 62
DB 252 GEDFLL--PPMVCVGCPIPVDSPLPEEALNHSIAKLNAEHDCTFYFKIDTVKKATVQV 309
QY 63 AGKYFIDFVARETTCKSKESNEELTSCETKLGQSLDCNAEVYVVPWEKKIYPTV 118
DB 310 GGLKYSIVFIARETTCKSGSNEELTKSCEINHGQILHCDANVYVVPWEKKYPTV 365
RESULT 7
A28055
K-kininogen, LMW I precursor - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 20-Jun-1989 #sequence_revision 20-Jun-1989 #text_change 15-Nov-1996
C:Accession: A28055
R:Furuto-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.
J. Biol. Chem. 260, 12054-12059, 1985
A:Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and
inhibitor.
A:Reference number: A92496; MUID:86008264; PMID:2413018
A:Accession: A28055
A:Molecule type: mRNA
A:Residues: 1-433 <FUR>
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-433/Product: K-kininogen, LMW I #status predicted <MAT>
F:19-131/Domain: cystatin homology <CY1>
F:142-253/Domain: cystatin homology <CY2>
F:264-375/Domain: cystatin homology <CY3>
Query Match 59.7%; Score 410; DB 2; Length 433;
Best Local Similarity 66.4%; Pred. No. 1.1e-30;
Matches 77; Conservative 13; Mismatches 26; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNATFYFKIDNVKKARVQV 62
DB 253 GDDLPELLPDCGCPNPVDSPLKEALGHSIAQNAENHTFYFKIDTVKKATSQVV 312
QY 63 AGKYFIDFVARETTCKSKESNEELTSCETKLGQSLDCNAEVYVVPWEKKIYPTV 118
DB 313 AGTYVIERFIARETTCKSKESNAELTADCTKLGQSLNCNANVYMPWENKVVPTV 368
RESULT 8
A25486
kininogen, HMW I precursor - rat
N:Contains: bradykinin
C:Species: Rattus norvegicus (Norway rat)
C:Date: 08-Mar-1989 #sequence_revision 08-Mar-1989 #text_change 15-Nov-1996
R:Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 262, 2190-2198, 1987
A:Title: Differing expression patterns and evolution of the rat kininogen gene family.
A:Reference number: A92625; MUID:87137443; PMID:3029068
A:Accession: A25486
A:Molecule type: mRNA
A:Residues: 1-639 <KIT>
A:Note: the authors translated the codon CAA for residue 347 as Asn
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-639/Product: kininogen, HMW I #status predicted <MAT>
F:19-131/Domain: cystatin homology <CY1>
F:142-253/Domain: cystatin homology <CY2>
F:264-375/Domain: cystatin homology <CY3>
Query Match 59.7%; Score 410; DB 2; Length 639;
Best Local Similarity 66.4%; Pred. No. 1.6e-30;
Matches 77; Conservative 13; Mismatches 26; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNATFYFKIDNVKKARVQV 62

F;Furuto-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.
J. Biol. Chem. 260, 12054-12059, 1985
A;Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and inhibitor.
A;Reference number: A92496; MUID:86008264; PMID:2413018
A;Accession: A01436
A;Molecule type: mRNA
A;Residues: 1-430 <FUR>
A;Cross-references: GB:M11883; NID:G205084; PIDN:AAA41489.1; PID:G205085
R;Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 262, 2190-2198, 1987
A;Title: Differing expression patterns and evolution of the rat kininogen gene family.
A;Reference number: A92625; MUID:97137443; PMID:3029068
A;Accession: D25486
A;Molecule type: DNA
A;Residues: 375-430 <KIT>
R;Enjyoji, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 973-979, 1988
A;Title: Purification and characterization of rat T-kininogens isolated from plasma of a rabbit.
A;Reference number: A92729; MUID:88087226; PMID:3121623
A;Accession: A28526
A;Molecule type: protein
A;Residues: 'E', 20-48; 376-430 <ENJ>
R;Kanda, S.; Sugiyama, K.; Takahashi, M.; Shumiya, S.; Tomino, S.; Nagase, S.
Jpn. J. Cancer Res. 81, 63-68, 1990
A;Title: Identification of a protein increasing in serum of Nagase analbuminemic rats by immunoblotting.
A;Reference number: P0193; MUID:90216390; PMID:2108948
A;Accession: P0193
A;Molecule type: mRNA
A;Residues: 330-420, 'R', 422-429, 'P' <KAN>
R;Anderson, K.P.; Croyle, M.L.; Lingrel, J.B.
Gene 81, 119-128, 1989
A;Title: Primary structure of a gene encoding rat T-kininogen.
A;Reference number: JQ0027; MUID:90034172; PMID:2806908
A;Accession: JQ0027
A;Molecule type: DNA
A;Residues: 1-60, 'E', 62-113, 'R', 115-165, 'F', 167-178, 'TKI', 182-211, 'P', 213-256, 'S', 258-386
R;Kageyama, R.; Kitamura, N.; Ohkubo, H.; Nakanishi, S.
J. Biol. Chem. 262, 2345-2351, 1987
A;Title: Differing utilization of homologous transcription initiation sites of rat K and I kininogen.
A;Reference number: A25488; MUID:87137465; PMID:3818598
A;Accession: B25488
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-48 <KAG>
A;Cross-references: GB:M14356; NID:G205090; PIDN:AAA41492.1; PID:G205091
R;Enjyoji, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 965-972, 1988
A;Title: Purification and characterization of two kinds of low molecular weight kininogen.
A;Reference number: A28525; MUID:88087225; PMID:3335530
A;Accession: A28525
A;Molecule type: protein
A;Residues: 376-430 <EN2>
R;Siererra, F.; Walter, R.; Vautravers, P.; Guigoz, Y.
Arch. Biochem. Biophys. 322, 333-338, 1995
A;Title: Identification of several isoforms of T-kininogen expressed in the liver of aged rats.
A;Reference number: S68034; MUID:96032652; PMID:7574705
A;Accession: S68036
A;Molecule type: mRNA
A;Residues: 340-430 <SIE>
A;Experimental source: clone pSG17
C;Comment: At least three types of LMW kininogen precursors are present in rat plasma, but only one is released after treatment with bradykinin.
C;Comment: T-kininogens contain T-kinin (I-S-bradykinin), a novel kinin isolated after treatment with bradykinin.
C;Comment: The T-kininogens are produced in response to an inflammatory stimulant.
C;Genetics: 65/3, 102/3, 130/1, 187/3, 223/2, 252/1, 309/3, 345/3, 374/3, 398/3
A;Introns: 65/3, 102/3, 130/1, 187/3, 223/2, 252/1, 309/3, 345/3, 374/3, 398/3
C;Superfamily: kininogen; cystatin homology
C;Keywords: acute phase; bradykinin; cysteine proteinase inhibitor; duplication; glycoprotein
F;1-18/Domain: signal sequence; #status predicted <SIG>
F;19-430/Product: T-kininogen I #status experimental <NAT>

F;19-130/Domain: cystatin homology <CV1>
F;141-252/Domain: cystatin homology <CV2>
F;263-374/Domain: cystatin homology <CV3>
F;378-386/Product: bradykinin #status predicted <BDY>
F;19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status experiment
F;82,126,168,204,326/Binding site: carboxylate (Asn) (covalent) #status predicted
F;83-94,107-125,141-144,205-217,228-247,263-266,327-339,350-369/disulfide bonds: #status
Query Match 55.3%; Score 380; DB 1; Length 430;
Best Local Similarity 62.1%; Pred. No. 6,7e-28;
Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;
QY 3 GKDFVQPTTKICVGGPRDIPNTPSELTEHTITPKLAENNATFYFKIDNVKARQVW 62
DB 252 GDDLFELLPKNCRGCPREIPVDSPELKEALGHSIAQLNAQNHIFYFKIDTVKKATQVW 311
QY 63 AGKYFIDFVARETTCSENEELTESCTKLGSLDCNAEYVVPVEKKIYPTV 118
DB 312 AGVIVIEFIARETNCQSKTELTADCTRHGLQSLNCNANVMPKENVKVPV 367
RESULT 13
UDSO
N;Alternate names: thiol proteinase inhibitor
C;Species: Bos primigenius taurus (cattle)
C;Date: 28-Feb-1986 #sequence_revision 28-Feb-1986 #text_change 06-Dec-1996
C;Accession: A01271
R;Hirado, M.; Tsunawaka, S.; Sakiyama, F.; Niinobe, M.; Fujii, S.
FEBS Lett. 186, 41-45, 1985
A;Title: Complete amino acid sequence of bovine colostrum low-M-r cysteine proteinase inhibitor.
A;Reference number: A01271; MUID:85231205; PMID:3891407
A;Accession: A01271
A;Molecule type: protein
A;Residues: 1-112 <HIR>
C;Superfamily: cystatin; cystatin homology
C;Keywords: colostrum; cysteine proteinase inhibitor
F;2-112/Domain: cystatin homology <CVS>
F;48-52/Region: inhibitory #status predicted
F;66-76,90-110/disulfide bonds: #status predicted
Query Match 20.0%; Score 137.5; DB 1; Length 112;
Best Local Similarity 28.8%; Pred. No. 7.5e-06;
Matches 32; Conservative 25; Mismatches 35; Indels 19; Gaps 4;
QY 24 NSPELEETHTITPKLAENNATFYFKIDNVKARQVVGAKYFIDFVARETTCSEKN 83
DB 12 NEEGVQELSAFSAVSEFNKSNDAVQSRVVRVVRARQVVGSMYFLDVELGRITCTK--S 69
QY 84 EELTESC-----ETKLGSLDCNAEYVVPVEKKIYPTVTVNWECE 126
DB 70 QANLDCPPHNQPHLAREKL-----CSFQYVVPVWNN-----TINLVKFSQ 111
RESULT 14
UDHU
N;Alternate names [validated] - human
C;Species: Homo sapiens (man)
C;Date: 06-Jul-1982 #sequence_revision 31-Mar-1991 #text_change 08-Dec-2000
C;Accession: S10216; S00004; J10095; A33400; S02751; A01270; A25434; S12288; A32732; A60
R;Abrahamson, M.; Olafsson, I.; Palsdottir, A.; Ulvbaeck, M.; Lundwall, A.; Jenson, O.
Biochem. J. 268, 287-294, 1990
A;Title: Structure and expression of the human cystatin C gene.
A;Reference number: S10216; MUID:90303202; PMID:2363674
A;Accession: S10216
A;Molecule type: DNA
A;Residues: 1-146 <AB1>
A;Cross-references: EMBL:X52255; NID:G30257; PIDN:CAA36497.1; PID:G296643
R;Abrahamson, M.; Grubb, A.; Olafsson, I.; Lundwall, A.
FEBS Lett. 216, 229-233, 1987
A;Title: Molecular cloning and sequence analysis of cDNA coding for the precursor of the
A;Reference number: S00004; MUID:87219149; PMID:3495457

A;Residues: 8-49 <ESN>
R;Esnard, A.; Esnard, F.; Guillou, F.; Gauthier, F.
FEBS Lett. 300, 131-135, 1992
A;Title: Production of the cysteine proteinase inhibitor cystatin C by rat Sertoli cells
A;Reference number: S21109; MUID:92225121; PMID:1563513
A;Accession: S21109
A;Molecule type: protein
A;Residues: 8, XX, 11-20 <ES2>
C;Superfamily: cystatin; cystatin homology
C;Keywords: cysteine proteinase inhibitor
F;16-127/Domain: cystatin homology <CYS>
F;80-90,104-124/Disulfide bonds: #status predicted

Query Match 18.9%; Score 130; DB 2; Length 127;
Best Local Similarity 28.0%; Pred. No. 4.3e-05;
Matches 30; Conservative 28; Mismatches 43; Indels 6; Gaps 4;

Qy 8 QPPTKICVGCPRDIPITNSPELEETLTHITIKLNANNATFYKIDNVKKARVQVAGKKY 67
Db 11 RPPRL-LGAPQADAGEEGVQRALDFAVSEYNKGSNDAYHSRAIQVVRARKQLVAGINY 69

Qy 68 PIDFVARETTCKESNEBELTESC---ETKLGQSLDCNAEYVVPWE 111
Db 70 YLDVEMGRITCTK-SQTNLT-NCPFHDQPHLMRKALCSFQIYSVPWK 114

Search completed: September 24, 2004, 14:10:49
Job time : 14.716 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:08:41 ; Search time 44.704 Seconds
(without alignments)
913.519 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687
Sequence: 1 GSKDFQVPPTKICVGCPRD.....VPWEKKIYPTVTVNHCECF 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1349238 seqs, 321558718 residues

Total number of hits satisfying chosen parameters: 1349238

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA:

1: /cgn2_6/ptodata/1/pubpaa/US07_PUBCOMB.pep.*
2: /cgn2_6/ptodata/1/pubpaa/PCT_NEW_PUB.pep.*
3: /cgn2_6/ptodata/1/pubpaa/US06_NEW_PUB.pep.*
4: /cgn2_6/ptodata/1/pubpaa/US06_PUBCOMB.pep.*
5: /cgn2_6/ptodata/1/pubpaa/US07_NEW_PUB.pep.*
6: /cgn2_6/ptodata/1/pubpaa/ECTUS_PUBCOMB.pep.*
7: /cgn2_6/ptodata/1/pubpaa/US08_NEW_PUB.pep.*
8: /cgn2_6/ptodata/1/pubpaa/US08_PUBCOMB.pep.*
9: /cgn2_6/ptodata/1/pubpaa/US09A_PUBCOMB.pep.*
10: /cgn2_6/ptodata/1/pubpaa/US09B_PUBCOMB.pep.*
11: /cgn2_6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubpaa/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	618	90.0	390	15	US-10-162-335-82
2	618	90.0	427	15	US-10-162-335-70
3	618	90.0	615	15	US-09-949-339-29
4	618	90.0	644	15	US-10-162-335-72
5	618	90.0	644	15	US-10-162-335-74
6	618	90.0	644	15	US-10-162-335-84
7	381	55.5	424	14	US-10-316-253-217
8	380	55.3	430	14	US-10-316-253-215
9	163.5	23.8	145	14	US-10-329-428-2
10	163.5	23.8	167	10	US-09-746-783-197
11	163	23.7	178	9	US-09-969-834-1
12	138.5	20.2	121	9	US-09-775-932-14
13	138.5	20.2	128	9	US-09-775-932-12
14	138.5	20.2	149	9	US-09-940-497-2
15	137.5	20.0	112	8	US-08-849-303-16

16	137.5	20.0	112	16	US-10-655-136-16
17	136.5	19.9	118	9	US-09-775-932-24
18	132.5	19.3	120	9	US-09-775-932-2
19	132.5	19.3	120	16	US-10-695-194-2
20	132.5	19.3	146	8	US-08-849-303-17
21	132.5	19.3	146	9	US-09-940-497-3
22	132.5	19.3	146	9	US-09-969-834-3
23	132.5	19.3	146	14	US-10-329-428-3
24	132.5	19.3	146	14	US-10-376-564-47
25	132.5	19.3	146	16	US-10-655-136-17
26	132.5	19.3	146	16	US-10-695-194-1
27	132.5	19.3	241	16	US-10-257-384A-2
28	132.5	19.3	641	16	US-10-257-384A-4
29	131.5	19.1	317	12	US-10-210-172-82
30	131.5	19.1	345	12	US-10-210-172-86
31	131.5	19.1	356	12	US-10-210-172-84
32	131.5	19.1	369	12	US-10-210-172-78
33	131.5	19.1	369	12	US-10-210-172-80
34	131.5	19.1	382	12	US-10-315-664-93
35	131.5	19.1	382	12	US-09-978-360A-425
36	130	18.9	127	8	US-08-849-303-19
37	130	18.9	127	16	US-10-655-136-19
38	129.5	18.9	140	14	US-10-376-564-46
39	129.5	18.9	140	14	US-10-376-564-48
40	128	18.6	111	8	US-08-849-303-26
41	128	18.6	111	16	US-10-655-136-26
42	127.5	18.6	140	8	US-08-849-303-18
43	127.5	18.6	140	16	US-10-655-136-18
44	124.5	18.1	121	9	US-09-775-932-8
45	124.5	18.1	141	8	US-08-849-303-24

ALIGNMENTS

RESULT 1

US-10-162-335-82

; Sequence 82, Application US/10162335

; Publication No. US20040009480A1

; GENERAL INFORMATION:

; APPLICANT: Anderson, David W.

; APPLICANT: Baumgartner, Jason C.

; APPLICANT: Boldog, Ferenc L.

; APPLICANT: Casman, Stacie J.

; APPLICANT: Edinger, Shlomit R.

; APPLICANT: Gangolli, Esha A.

; APPLICANT: Gerlach, Valerie

; APPLICANT: Gorman, Linda

; APPLICANT: Guo, Xiaojia (Sasha)

; APPLICANT: Hjalte, Tord

; APPLICANT: Kekuda, Ramesh

; APPLICANT: Li, Li

; APPLICANT: MacDougall, John R.

; APPLICANT: Malyankar, Uriel M.

; APPLICANT: Miller, Isabelle

; APPLICANT: Padigaru, Muralidhara

; APPLICANT: Patturajan, Meera

; APPLICANT: Pena, Carol E. A.

; APPLICANT: Rastelli, Luca

; APPLICANT: Shimkets, Richard A.

; APPLICANT: Stone, David J.

; APPLICANT: Spytek, Kimberly A.

; APPLICANT: Vernet, Corine A. M.

; APPLICANT: Voss, Edward Z.

; APPLICANT: Zerhusen, Bryan D.

; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method

; FILE REFERENCE: 21402-377 B

; CURRENT APPLICATION NUMBER: US/10162,335

; CURRENT FILING DATE: 2002-10-01

; PRIOR APPLICATION NUMBER: 60/295,607

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/295,661

; PRIOR FILING DATE: 2001-06-04

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; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 82
; LENGTH: 390
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-82

Query Match          90.0%; Score 618; DB 15; Length 390;
Best Local Similarity 100.0%; Pred. No. 5.7e-59;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDVPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 62
Db 216 GKDVPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 275

QY 63 AGKGYFIDFVARETTCSKESNEELTESCETKKLGQSLDCNAEYVVPWEKKIYPTV 118
Db 276 AGKGYFIDFVARETTCSKESNEELTESCETKKLGQSLDCNAEYVVPWEKKIYPTV 331

RESULT 2
US-10-162-335-70
; Sequence 70, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Salomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjal, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigar, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shimkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; APPLICANT: Zernusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
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; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 70
; LENGTH: 398
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-70

Query Match          90.0%; Score 618; DB 15; Length 398;
Best Local Similarity 100.0%; Pred. No. 5.8e-59;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDVPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 62
Db 224 GKDVPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 283

QY 63 AGKGYFIDFVARETTCSKESNEELTESCETKKLGQSLDCNAEYVVPWEKKIYPTV 118
Db 284 AGKGYFIDFVARETTCSKESNEELTESCETKKLGQSLDCNAEYVVPWEKKIYPTV 339

RESULT 3
US-09-919-039-29
; Sequence 29, Application US/09919039
; Publication No. US20030108871A1
; GENERAL INFORMATION:
; APPLICANT: Kaser, Matthew R.
; TITLE OF INVENTION: GENES EXPRESSED IN TREATED HUMAN C3A LIVER CELL CULTURES
; FILE REFERENCE: PA-0035 US
; CURRENT APPLICATION NUMBER: US/09/919,039
; CURRENT FILING DATE: 2002-09-09
; PRIOR APPLICATION NUMBER: 60/222,113
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 401
; SOFTWARE: PERL Program
; SEQ ID NO 29
; LENGTH: 427
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: incyte ID No. US20030108871A1 167507CD1
US-09-919-039-29

Query Match          90.0%; Score 618; DB 10; Length 427;
Best Local Similarity 100.0%; Pred. No. 6.4e-59;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDVPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 62
Db 253 GKDVPPTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARQVW 312

QY 63 AGKGYFIDFVARETTCSKESNEELTESCETKKLGQSLDCNAEYVVPWEKKIYPTV 118
Db 313 AGKGYFIDFVARETTCSKESNEELTESCETKKLGQSLDCNAEYVVPWEKKIYPTV 368
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RESULT 4
US-10-162-335-72
; Sequence 72, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalte, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigaru, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shimkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; APPLICANT: Zerhusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 72
; LENGTH: 615
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-72
Query Match 90.0%; Score 618; DB 15; Length 615;
Best Local Similarity 100.0%; Pred. No. 1e-58;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDIPNTPSPELEETLTHITIKLAENNATFYFKIDNVKKARVQV 62
DB 224 GKDFVQPTKICVGCPRDIPNTPSPELEETLTHITIKLAENNATFYFKIDNVKKARVQV 283
QY 63 AGKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTV 118

DB 284 AGKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTV 339
RESULT 5
US-10-162-335-74
; Sequence 74, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalte, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigaru, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shimkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; APPLICANT: Zerhusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 74
; LENGTH: 644
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-74
Query Match 90.0%; Score 618; DB 15; Length 644;
Best Local Similarity 100.0%; Pred. No. 1.1e-58;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDIPNTPSPELEETLTHITIKLAENNATFYFKIDNVKKARVQV 62
DB 253 GKDFVQPTKICVGCPRDIPNTPSPELEETLTHITIKLAENNATFYFKIDNVKKARVQV 312

QY 63 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNNAEVVVPWEKKIYPTV 118
Db 313 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNNAEVVVPWEKKIYPTV 368

RESULT 6

US-10-162-335-84

; Sequence 84, Application US/10162335

; Publication No. US20040009480A1

; GENERAL INFORMATION:

; APPLICANT: Anderson, David W.

; APPLICANT: Baumgartner, Jason C.

; APPLICANT: Boldog, Ferenc L.

; APPLICANT: Casman, Stacie J.

; APPLICANT: Edinger, Shlomit R.

; APPLICANT: Gangolli, Esha A.

; APPLICANT: Gerlach, Valerie

; APPLICANT: Gorman, Linda

; APPLICANT: Guo, Xiaojia (Sasha)

; APPLICANT: Hjal, Tord

; APPLICANT: Kekuda, Ramesh

; APPLICANT: Li, Li

; APPLICANT: MacDougall, John R.

; APPLICANT: Malyankar, Uriel M.

; APPLICANT: Millet, Isabelle

; APPLICANT: Padigaru, Muralidhara

; APPLICANT: Patturajan, Meera

; APPLICANT: Pena, Carol E. A.

; APPLICANT: Rastelli, Luca

; APPLICANT: Shimkets, Richard A.

; APPLICANT: Stone, David J.

; APPLICANT: Spytek, Kimberly A.

; APPLICANT: Vernet, Corine A. M.

; APPLICANT: Voss, Edward Z.

; APPLICANT: Zerhusen, Bryan D.

; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method

; FILE REFERENCE: 21402-377 B

; CURRENT APPLICATION NUMBER: US/10/162,335

; CURRENT FILING DATE: 2002-10-01

; PRIOR APPLICATION NUMBER: 60/295,607

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/295,661

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/296,404

; PRIOR FILING DATE: 2001-06-06

; PRIOR APPLICATION NUMBER: 60/296,418

; PRIOR FILING DATE: 2001-06-06

; PRIOR APPLICATION NUMBER: 60/297,414

; PRIOR FILING DATE: 2001-06-11

; PRIOR APPLICATION NUMBER: 60/297,567

; PRIOR FILING DATE: 2001-06-12

; PRIOR APPLICATION NUMBER: 60/298,285

; PRIOR FILING DATE: 2001-06-14

; PRIOR APPLICATION NUMBER: 60/298,556

; PRIOR FILING DATE: 2001-05-15

; PRIOR APPLICATION NUMBER: 60/299,949

; PRIOR FILING DATE: 2001-06-21

; PRIOR APPLICATION NUMBER: 60/300,883

; PRIOR FILING DATE: 2001-06-26

; Remaining prior application data removed - See File Wrapper or PALM.

; SEQ ID NO 84

; LENGTH: 644

; TYPE: PRT

; ORGANISM: Homo sapiens

US-10-162-335-84

Query Match

Best Local Similarity 90.0%; Score 618; DB 15; Length 644;

Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNSPELEETLTHITKLNAENNAFFYFKIDNVKARQVV 62

Db 253 GKDFVQPTKICVGCPRDIPNSPELEETLTHITKLNAENNAFFYFKIDNVKARQVV 312
QY 63 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNNAEVVVPWEKKIYPTV 118
Db 313 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNNAEVVVPWEKKIYPTV 368

RESULT 7

US-10-316-253-217

; Sequence 217, Application US/10316253

; Publication No. US20030162706A1

; GENERAL INFORMATION:

; APPLICANT: The Procter & Gamble Company

; APPLICANT: Peters, Kevin

; APPLICANT: Thompson, Larry

; APPLICANT: Wang, Feng

; APPLICANT: Greis, Kenneth

; TITLE OF INVENTION: Angiogenesis Modulating Proteins

; FILE REFERENCE: 8865M

; CURRENT APPLICATION NUMBER: US/10/316,253

; CURRENT FILING DATE: 2002-12-10

; PRIOR APPLICATION NUMBER: US 60/355,295

; PRIOR FILING DATE: 2002-02-08

; NUMBER OF SEQ ID NOS: 308

; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 217

; LENGTH: 424

; TYPE: PRT

; ORGANISM: Rattus norvegicus

US-10-316-253-217

Query Match

Best Local Similarity 55.5%; Score 381; DB 14; Length 424;

Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNSPELEETLTHITKLNAENNAFFYFKIDNVKARQVV 62
Db 246 GDDLPELLPKNCRGCPREIPVDSPELKEALGHISARLNAQHIFYFKIDTVKATQVV 305

QY 63 AGKYFIDFVARETTCSKESNEELTESCETKKGQSLDCNNAEVVVPWEKKIYPTV 118
Db 306 AGVIYVIERIARETNCSKQSKTELTADCTKHLGQSLCNCNANVYMPWENKVVPTV 361

RESULT 8

US-10-316-253-215

; Sequence 215, Application US/10316253

; Publication No. US20030162706A1

; GENERAL INFORMATION:

; APPLICANT: The Procter & Gamble Company

; APPLICANT: Peters, Kevin

; APPLICANT: Thompson, Larry

; APPLICANT: Wang, Feng

; APPLICANT: Greis, Kenneth

; TITLE OF INVENTION: Angiogenesis Modulating Proteins

; FILE REFERENCE: 8865M

; CURRENT APPLICATION NUMBER: US/10/316,253

; CURRENT FILING DATE: 2002-12-10

; PRIOR APPLICATION NUMBER: US 60/355,295

; PRIOR FILING DATE: 2002-02-08

; NUMBER OF SEQ ID NOS: 308

; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 215

; LENGTH: 430

; TYPE: PRT

; ORGANISM: Rattus norvegicus

US-10-316-253-215

Query Match

Best Local Similarity 55.3%; Score 380; DB 14; Length 430;

Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;

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QY 3 GKDFVQPTKICVCPDIPNTSPLEETLTHITKLNNAENATFYFKIDNVKARQVV 62
DB 252 GDDLPELLPKNCRCFPIPDSPELKEALGHSIAQLNAQHNIIFYKIDTVKATQVV 311
QY 63 AGKYFIDFVARETTCKSNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTV 118
DB 312 AGVYVIEFTARETNCQSKSTELTADCTKHLGSLNCNANVTYMPWENKVVPTV 367

RESULT 9
US-10-329-428-2
; Sequence 2, Application US/10329428
; Publication No. US20030114646A1
; GENERAL INFORMATION:
; APPLICANT: Li, et al.
; TITLE OF INVENTION: Human Cystatin F
; FILE REFERENCE: P2265P1D2
; CURRENT APPLICATION NUMBER: US/10/329,428
; CURRENT FILING DATE: 2002-12-27
; PRIOR APPLICATION NUMBER: 60/014,795
; PRIOR FILING DATE: 1996-04-03
; PRIOR APPLICATION NUMBER: 08/832,535
; PRIOR FILING DATE: 1997-04-03
; PRIOR APPLICATION NUMBER: 09/019,485
; PRIOR FILING DATE: 1998-01-29
; PRIOR APPLICATION NUMBER: 09/528,436
; PRIOR FILING DATE: 2000-03-17
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: Patent in version 3.2
; SEQ ID NO 2
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-329-428-2

Query Match 23.8%; Score 163.5; DB 14; Length 145;
Best Local Similarity 31.6%; Pred No. 9.7e-10;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVCPDIPNTSPLEETLTHITKLNNAENATFYFKIDNVKARQVVAGKYFID 70
DB 32 SRVKGPFKTIKNDPGVLQAARYSVKFNCTNDMLFKESRITRALVQIVKGLKYLE 91
QY 71 FVARETTCKSNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVTVAHWE 124
DB 92 VEIGRTCKKQHLRL-DDCDFQNTLTKQLTSLCYSEVVWVPW-----LQHF 138

RESULT 10
US-09-746-783-197
; Sequence 197, Application US/09746783
; Publication No. US20030044935A1
; GENERAL INFORMATION:
; APPLICANT: Jacobs, Kenneth
; McCoy, John M.
; LaVallie, Edward R.
; Racie, Lisa A.
; Treacy, Maurice
; Spaulding, Vikki
; Agostino, Michael J.
; Howes, Steven H.
; Fechtel, Kim
; TITLE OF INVENTION: SECRETED PROTEINS AND POLYNUCLEOTIDES
; ENCODING THEM
; NUMBER OF SEQUENCES: 231
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genetics Institute, Inc.
; STREET: 87 CambridgePark Drive
; CITY: Cambridge
; STATE: MA
; COUNTRY: U.S.A.
; ZIP: 02140
; COMPUTER READABLE FORM:

QY 3 GKDFVQPTKICVCPDIPNTSPLEETLTHITKLNNAENATFYFKIDNVKARQVV 62
DB 252 GDDLPELLPKNCRCFPIPDSPELKEALGHSIAQLNAQHNIIFYKIDTVKATQVV 311
QY 63 AGKYFIDFVARETTCKSNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTV 118
DB 312 AGVYVIEFTARETNCQSKSTELTADCTKHLGSLNCNANVTYMPWENKVVPTV 367

RESULT 11
US-09-969-834-1
; Sequence 1, Application US/09969834
; Patent No. US20020102711A1
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; Goli, Surya K.
; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE
; PROTEIN
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/969,834
; FILING DATE: 01-Oct-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/09/471,765
; FILING DATE: <Unknown>
; APPLICATION NUMBER: US/08/791,522
; FILING DATE: <Unknown>
; APPLICATION NUMBER: 09/471,765
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
```

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; REFERENCE/DOCKET NUMBER: PF-0193 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 178 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: 30443
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-969-834-1

Query Match      23.7%; Score 163; DB 9; Length 178;
Best Local Similarity 34.0%; Pred. No. 1.4e-09;
Matches 35; Conservative 20; Mismatches 44; Indels 4; Gaps 2;

QY 11 TKICVGCPRDIPNTPSPELEETLTHITIKLNAENNAFYFKIDNVKKARVQVVGKKYFI 70
DB 54 SRVKGFPKTIKNDGVIQAARISVEKENNCTNMFLEKSRITRALVQIVKGLKYMIE 113
QY 71 FVARETTCSKE---TKKLGQSLDCNAEVVVPWE 110
DB 114 VEIGRTTCKKNQHLRL-DCDDFQTNHTLAKQTLSCYSEVVVVPW 155

RESULT 12
US-09-775-932-14
; Sequence 14, Application US/09775932
; Patent No. US20020137671A1
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent in Ver. 2.0
; SEQ ID NO 14
; LENGTH: 121
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-14

Query Match      20.2%; Score 138.5; DB 9; Length 121;
Best Local Similarity 31.5%; Pred. No. 4.2e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPNTPSPELEETLTHITIKLNAENNAFYFKIDNVKKARVQVVGKKYFI 69
DB 2 PQERWVGELRDSPPDPQVQKAAQAAVSYNGMSNIYFRDTHIIKAQSQLVAGIKYFL 61
QY 70 DFVARETTCSKE---SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
DB 62 TMEMGSTDCRKTTRVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 108

RESULT 13
US-09-775-932-12
; Sequence 12, Application US/09775932
; Patent No. US20020137671A1
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02

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```

; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: Patent in Ver. 2.0
; SEQ ID NO 12
; LENGTH: 128
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-12

Query Match      20.2%; Score 138.5; DB 9; Length 128;
Best Local Similarity 31.5%; Pred. No. 4.5e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPNTPSPELEETLTHITIKLNAENNAFYFKIDNVKKARVQVVGKKYFI 69
DB 9 PQERWVGELRDSPPDPQVQKAAQAAVSYNGMSNIYFRDTHIIKAQSQLVAGIKYFL 68
QY 70 DFVARETTCSKE---SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
DB 69 TMEMGSTDCRKTTRVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 115

RESULT 14
US-09-940-497-2
; Sequence 2, Application US/09940497
; Patent No. US20020052476A1
; GENERAL INFORMATION:
; APPLICANT: Ni et al.
; TITLE OF INVENTION: Human Cystatin E
; CURRENT APPLICATION NUMBER: US/09/940,497
; CURRENT FILING DATE: 2001-08-29
; PRIOR APPLICATION NUMBER: US 09/241,376
; PRIOR FILING DATE: 1999-02-02
; PRIOR APPLICATION NUMBER: US 08/744,138
; PRIOR FILING DATE: 1996-11-05
; PRIOR APPLICATION NUMBER: US 08/461,030
; PRIOR FILING DATE: 1995-06-05
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 2
; LENGTH: 149
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-940-497-2

Query Match      20.2%; Score 138.5; DB 9; Length 149;
Best Local Similarity 31.5%; Pred. No. 5.5e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 10 PTKICVGCPRDIPNTPSPELEETLTHITIKLNAENNAFYFKIDNVKKARVQVVGKKYFI 69
DB 30 PQERWVGELRDSPPDPQVQKAAQAAVSYNGMSNIYFRDTHIIKAQSQLVAGIKYFL 89
QY 70 DFVARETTCSKE---SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
DB 90 TMEMGSTDCRKTTRVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 136

RESULT 15
US-08-849-303-16
; Sequence 16, Application US/08849303
; Publication No. US20030221209A1
; GENERAL INFORMATION:
; APPLICANT: Atkinson, Howard J.
; APPLICANT: McPherson, Michael J.
; APPLICANT: Urwin, Peter E.
; TITLE OF INVENTION: MODIFIED PROTEINASE INHIBITORS
; NUMBER OF SEQUENCES: 79
; CORRESPONDENCE ADDRESS:

```


ADDRESSEE: Klauber & Jackson
STREET: 411 Hackensack Avenue, 4th Floor
CITY: Hackensack
STATE: New Jersey
COUNTRY: USA
ZIP: 07601
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/849,303
FILING DATE: 21-MAY-1997
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Jackson Esq., David A.
REGISTRATION NUMBER: 26,742
REFERENCE/DOCKET NUMBER: 1321-1-003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 201-487-5800
TELEFAX: 201-343-1684
TELEX: 133521
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 112 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
US-08-849-303-16

Query Match 20.0%; Score 137.5; DB 8; Length 112;
Best Local Similarity 28.8%; Pred. No. 4.9e-07;
Matches 32; Conservative 25; Mismatches 35; Indels 19; Gaps 4;
Qy 24 NSPELEETLTHTITKLNANNATYFKIDNVKARVGVAGKYFIDFVARETTCSKSN 83
Db 12 NEEGVQELSPAVSEFNKRSNDAYQSRVVRVRAKQVVGCMYFLDVELGRITCTK--S 69
Qy 84 BELTSC-----ETKKLQGLDCNAEVVVPWEKKIYPTVTVNHWECE 126
Db 70 QANLDCSPFHNPQLKREKL-----CSFQYVVPWMN---TINLVKFSQ 111

Search completed: September 24, 2004, 14:13:04
Job time : 45.704 secs

GenCore version 5.1.6
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QM protein - protein search, using sw model

Run on: September 24, 2004, 14:07:01 ; Search time 14.268 Seconds
(without alignments)
445.051 Million cell updates/sec

Title: US-10-661-784-1
Perfect score: 660
Sequence: 1 GKDFVQPTKICVGCPRDIP.....YVVPWEKKIYPTWNCQPLGM 123

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues
Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA.*
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3: /cgn2_6/ptodata/2/iaa/6A COMB.pcp.*
4: /cgn2_6/ptodata/2/iaa/6B COMB.pcp.*
5: /cgn2_6/ptodata/2/iaa/PCTUS COMB.pcp.*
6: /cgn2_6/ptodata/2/iaa/backfiles1.pcp.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	600	90.9	117	1	US-08-193-114B-1
2	594	90.0	117	5	PCT-US92-06809-1
3	169	25.6	178	2	US-08-791-522-1
4	169	25.6	178	3	US-09-314-777-1
5	165.5	25.1	145	2	US-08-832-535-2
6	165.5	25.1	145	3	US-09-019-485-2
7	165.5	25.1	145	3	US-09-019-485-3
8	165.5	25.1	145	3	US-09-431-480-9
9	165.5	25.1	145	3	US-09-617-302-9
10	165.5	25.1	145	4	US-09-528-436B-2
11	155	23.5	27	3	US-08-676-242-11
12	145	22.0	64	3	US-08-676-242-22
13	145	22.0	64	3	US-09-402-732-1
14	138.5	21.0	121	4	US-09-775-932-14
15	138.5	21.0	128	4	US-09-775-932-12
16	138.5	21.0	149	2	US-08-461-030C-2
17	138.5	21.0	149	3	US-08-744-138-2
18	138.5	21.0	149	3	US-09-431-480-8
19	138.5	21.0	149	3	US-09-431-480-10
20	138.5	21.0	149	3	US-09-617-302-8
21	138.5	21.0	149	3	US-09-617-302-10
22	138.5	21.0	149	4	US-09-241-376-2
23	138.5	21.0	149	4	US-09-940-497-2
24	136	20.6	112	4	US-08-849-303-16
25	135	20.5	118	4	US-09-775-932-24
26	134	20.3	148	5	PCT-US95-07135-2
27	132	20.0	26	3	US-08-676-242-15

28	130	19.7	127	4	US-08-849-303-19	Sequence 19, Appl
29	129.5	19.6	140	4	US-08-866-319A-46	Sequence 46, Appl
30	129.5	19.6	140	4	US-08-866-319A-48	Sequence 48, Appl
31	129.5	19.6	382	4	US-09-599-360B-93	Sequence 93, Appl
32	128.5	19.5	146	6	5432264-6	Patent No. 5432264
33	128	19.4	111	4	US-08-849-303-26	Sequence 26, Appl
34	127.5	19.3	120	4	US-09-775-932-2	Sequence 2, Appl
35	127.5	19.3	140	3	US-09-431-480-5	Sequence 5, Appl
36	127.5	19.3	140	3	US-08-617-302-5	Sequence 5, Appl
37	127.5	19.3	140	4	US-08-849-303-18	Sequence 18, Appl
38	127.5	19.3	145	2	US-08-832-535-11	Sequence 11, Appl
39	127.5	19.3	146	2	US-08-791-522-3	Sequence 3, Appl
40	127.5	19.3	146	3	US-08-744-138-3	Sequence 3, Appl
41	127.5	19.3	146	3	US-09-019-485-4	Sequence 4, Appl
42	127.5	19.3	146	3	US-09-314-777-3	Sequence 3, Appl
43	127.5	19.3	146	3	US-09-431-480-6	Sequence 6, Appl
44	127.5	19.3	146	3	US-09-617-302-6	Sequence 6, Appl
45	127.5	19.3	146	4	US-09-241-376-3	Sequence 3, Appl

ALIGNMENTS

RESULT 1
US-08-193-114B-1
; Sequence 1, Application US/08193114B
; Patent No. 5472945
; GENERAL INFORMATION:
; APPLICANT: Schmaier, Alvin H.
; APPLICANT: Jiang, Yongping
; TITLE OF INVENTION: Modulation of Blood
; TITLE OF INVENTION: Pressure and Inhibition of Platelet Activation
; TITLE OF INVENTION: with Kininogen Fragment
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Seidel, Gonda, Lavoragna &
; ADDRESSEE: Monaco, P.C.
; STREET: 1800 Two Penn Center Plaza
; CITY: Philadelphia
; STATE: Pennsylvania
; COUNTRY: U.S.A.
; ZIP: 19102

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/193,114B
FILING DATE: 9 February 1994
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: U.S. Application
FILING DATE: 13 August 1991
REFERENCE NUMBER: Serial No. 5472945 07/744,545
ATTORNEY/AGENT INFORMATION:
NAME: Monaco, Daniel A.
REGISTRATION NUMBER: 30,480
REFERENCE/DOCKET NUMBER: 6056-137 CII
TELECOMMUNICATION INFORMATION:
TELEPHONE: (215) 568-8383
TELEFAX: (215) 568-5549
TELEX: No. 5472345e
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 117 amino acids
TYPE: peptide
TOPOLOGY: linear

US-08-193-114B-1
Query Match 90.9%; Score 600; DB 1; Length 117;
Best Local Similarity 100.0%; Pred. No. 5.8e-59;
Matches 112; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 12 CVGCPDIPNTPSPLEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFIDFVA 71
 Db 1 CVGCPDIPNTPSPLEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFIDFVA 60
 QY 72 RETTCSKESNEBELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 123
 Db 61 RETTCSKESNEBELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 112

RESULT 2

PCT-US92-06809-1

; Sequence 1, Application PC/TUS9206809

; GENERAL INFORMATION:

; APPLICANT: Schmaier, Alvin H.

; APPLICANT: Jiang, Yongping

; TITLE OF INVENTION: Modulation of Blood

; TITLE OF INVENTION: Pressure by Altering Bradykinin Levels

; NUMBER OF SEQUENCES: 2

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Temple University - Of the

; ADDRESSEE: Commonwealth System of Higher Education

; STREET: 406 University Services

; CITY: Philadelphia

; STATE: Pennsylvania

; COUNTRY: U.S.A.

; ZIP: 19122

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb

; COMPUTER: IBM PS/2

; OPERATING SYSTEM: MS-DOS

; SOFTWARE: WordPerfect 5.1

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: PCT/US92/06809

; FILING DATE: 19910813

; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: U.S. Application

; APPLICATION NUMBER: Serial No. 744,545

; FILING DATE: 13 August 1991

; ATTORNEY/AGENT INFORMATION:

; NAME: Monaco, Daniel A.

; REGISTRATION NUMBER: 30,480

; REFERENCE/DOCKET NUMBER: 6056-137

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (215) 568-8383

; TELEFAX: (215) 568-5549

; TELEX:

; INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 117 amino acids

; TYPE: AMINO ACID

; TOPOLOGY: linear

PCT-US92-06809-1

Query Match

Best Local Similarity 90.0%; Score 594; DB 5; Length 117;

Matches 110; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 12 CVGCPDIPNTPSPLEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFIDFVA 71

Db 1 CVGCPDIPNTPSPLEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFIDFVA 60

QY 72 RETTCSKESNEBELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 123

Db 61 RETTCSKESNEBELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 112

RESULT 3

US-08-791-522-1

; Sequence 1, Application US/08791522

; Patent No. 5935817

; GENERAL INFORMATION:
 ; APPLICANT: Bandman, Olga
 ; APPLICANT: Goli, Surya K.
 ; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE
 ; TITLE OF INVENTION: PROTEIN
 ; NUMBER OF SEQUENCES: 4
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Incyte Pharmaceuticals, Inc.
 ; STREET: 3174 Porter Drive
 ; CITY: Palo Alto
 ; STATE: CA
 ; COUNTRY: USA
 ; ZIP: 94304

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Diskette

; COMPUTER: IBM Compatible

; OPERATING SYSTEM: DOS

; SOFTWARE: FastSeq for Windows Version 2.0

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/791,522

; FILING DATE: Filed Herewith

; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER:

; FILING DATE:

; ATTORNEY/AGENT INFORMATION:

; NAME: Billings, Lucy J.

; REGISTRATION/DOCKET NUMBER: 36,749

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 415-855-0555

; TELEFAX: 415-845-4166

; INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 178 amino acids

; TYPE: amino acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; IMMEDIATE SOURCE:

; CLONE: 30443

US-08-791-522-1

Query Match

Best Local Similarity 25.6%; Score 169; DB 2; Length 178;

Matches 39; Conservative 22; Mismatches 49; Indels 10; Gaps 4;

QY 9 TKICVCPDIPNTPSPLEETLTHITIKLNAENNAFFYFKIDNVKARVQVWAGKYFID 68

Db 54 SRVKGPFKTIKNDPGLQAARYSVEKFNCTNDMLFKESRITRALVQIVKGLKYLE 113

QY 69 FVARETTCSKESNEBELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 123

Db 114 VEIGRTCKKQHLRL-DCCDFQTNHTLKTLCYSEVWVVPW-----VPALRGAGCSPLSL 168

RESULT 4

US-09-314-777-1

; Sequence 1, Application US/09314777

; Patent No. 6110686

; GENERAL INFORMATION:

; APPLICANT: Bandman, Olga

; APPLICANT: Goli, Surya K.

; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE

; TITLE OF INVENTION: PROTEIN

; NUMBER OF SEQUENCES: 4

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Incyte Pharmaceuticals, Inc.

; STREET: 3174 Porter Drive

; CITY: Palo Alto

; STATE: CA

; COUNTRY: USA

; ZIP: 94304

; COMPUTER READABLE FORM:

;; PRIOR FILING DATE: 1998-01-29
;; PRIOR APPLICATION NUMBER: 08/832,555
;; PRIOR FILING DATE: 1999-04-03
;; PRIOR APPLICATION NUMBER: 60/014,795
;; PRIOR FILING DATE: 1996-04-03
;; NUMBER OF SEQ ID NOS: 16
;; SOFTWARE: Patent in version 3.2
;; SEQ ID NO 2
;; LENGTH: 145
;; TYPE: PRT
;; ORGANISM: Homo sapiens
US-09-528-436B-2
Query Match 25.1%; Score 165.5; DB 4; Length 145;
Best Local Similarity 32.5%; Pred. No. 1e-10;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;
Qy 9 TKICVGPDRPTNSPELEETLTTITIKLNAENATYFKIDNVKARQVAVGKYYFID 68
Db 32 SRVKGPFPTIKTNDPGVLQAARYSVEKFNNTNDMFLKESRITRALVOIVKGLKYMLE 91
Qy 69 FVARETTCKESNEELTESCS--TKLQSLDCNAEVVVPWEKKI-YPTVNC 118
Db 92 VEIGRTCKCKQHRL-LDCCDFQTNHILKQTLSCYSEVVVPWLQHPVPVLR 144
RESULT 11
US-08-676-242-11
;; Sequence 11, Application US/08676242C
;; Patent No. 6143719
;; GENERAL INFORMATION:
;; APPLICANT: The Regents of the University of Michigan
;; APPLICANT: Schmaier, Alvin H.
;; APPLICANT: Hasan, Ahmed A.K.
;; TITLE OF INVENTION: Bradykinin Analogs As Selective Thrombin Inhibitors
;; FILE REFERENCE: 8820-2 US
;; CURRENT APPLICATION NUMBER: US/08/676,242C
;; PRIOR FILING DATE: 2000-07-16
;; EARLIER APPLICATION NUMBER: 60/000,096
;; EARLIER FILING DATE: 1995-06-09
;; PRIOR FILING DATE: 1996-06-07
;; NUMBER OF SEQ ID NOS: 24
;; SOFTWARE: Patent in Ver. 2.1
;; SEQ ID NO 11
;; LENGTH: 27
;; TYPE: PRT
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: Description of Artificial Sequence: Bradykinin
US-08-676-242-11
Query Match 23.5%; Score 155; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 1.6e-10;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 97 LDCNAEVVVPWEKKIYPTVNCQPLGM 123
Db 1 LDCNAEVVVPWEKKIYPTVNCQPLGM 27
RESULT 12
US-08-676-242-22
;; Sequence 22, Application US/08676242C
;; Patent No. 6143719
;; GENERAL INFORMATION:
;; APPLICANT: The Regents of the University of Michigan
;; APPLICANT: Schmaier, Alvin H.
;; APPLICANT: Hasan, Ahmed A.K.
;; TITLE OF INVENTION: Bradykinin Analogs As Selective Thrombin Inhibitors
;; FILE REFERENCE: 8820-2 US
;; CURRENT APPLICATION NUMBER: US/08/676,242C
US-09-402-732-1
;; Sequence 1, Application US/09402732
;; Patent No. 6251855
;; GENERAL INFORMATION:
;; APPLICANT: Schmaier, Alvin H.
;; APPLICANT: Hasan, A.K. Ahmed
;; TITLE OF INVENTION: Bradykinin Analogs As Selective Inhibitors of Cell
;; TITLE OF INVENTION: Activation
;; FILE REFERENCE: 8820-3
;; CURRENT APPLICATION NUMBER: US/09/402,732
;; CURRENT FILING DATE: 1999-12-01
;; PRIOR APPLICATION NUMBER: 60/046,085
;; PRIOR FILING DATE: 1997-04-23
;; PRIOR APPLICATION NUMBER: PCT/US98/08015
;; PRIOR FILING DATE: 1998-04-21
;; NUMBER OF SEQ ID NOS: 10
;; SOFTWARE: Patent in Ver. 2.1
;; SEQ ID NO 1
;; LENGTH: 64
;; TYPE: PRT
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: Description of Artificial Sequence: Segment of
;; OTHER INFORMATION: human kininogen (residues 333-396 thereof)
US-09-402-732-1
Query Match 22.0%; Score 145; DB 3; Length 64;
Best Local Similarity 100.0%; Pred. No. 6.3e-09;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 99 CNAEVVVPWEKKIYPTVNCQPLGM 123
Db 1 CNAEVVVPWEKKIYPTVNCQPLGM 25
RESULT 13
US-09-402-732-1
;; Sequence 1, Application US/09402732
;; Patent No. 6251855
;; GENERAL INFORMATION:
;; APPLICANT: Schmaier, Alvin H.
;; APPLICANT: Hasan, A.K. Ahmed
;; TITLE OF INVENTION: Bradykinin Analogs As Selective Inhibitors of Cell
;; TITLE OF INVENTION: Activation
;; FILE REFERENCE: 8820-3
;; CURRENT APPLICATION NUMBER: US/09/402,732
;; CURRENT FILING DATE: 1999-12-01
;; PRIOR APPLICATION NUMBER: 60/046,085
;; PRIOR FILING DATE: 1997-04-23
;; PRIOR APPLICATION NUMBER: PCT/US98/08015
;; PRIOR FILING DATE: 1998-04-21
;; NUMBER OF SEQ ID NOS: 10
;; SOFTWARE: Patent in Ver. 2.1
;; SEQ ID NO 1
;; LENGTH: 64
;; TYPE: PRT
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: Description of Artificial Sequence: Segment of
;; OTHER INFORMATION: human kininogen (residues 333-396 thereof)
US-09-402-732-1
Query Match 22.0%; Score 145; DB 3; Length 64;
Best Local Similarity 100.0%; Pred. No. 6.3e-09;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 99 CNAEVVVPWEKKIYPTVNCQPLGM 123
Db 1 CNAEVVVPWEKKIYPTVNCQPLGM 25
RESULT 14
US-09-775-932-14
;; Sequence 14, Application US/09775932
;; Patent No. 6534477
;; GENERAL INFORMATION:
;; APPLICANT: University of British Columbia
;; TITLE OF INVENTION: Production and use of Modified Cystatins
;; FILE REFERENCE: 58069
;; CURRENT APPLICATION NUMBER: US/09/775,932
;; CURRENT FILING DATE: 2001-02-02
;; PRIOR APPLICATION NUMBER: CA99/00717
;; PRIOR FILING DATE: 1999-08-05

;; PRIOR FILING DATE: 1998-01-29
;; PRIOR APPLICATION NUMBER: 08/832,555
;; PRIOR FILING DATE: 1999-04-03
;; PRIOR APPLICATION NUMBER: 60/014,795
;; PRIOR FILING DATE: 1996-04-03
;; NUMBER OF SEQ ID NOS: 16
;; SOFTWARE: Patent in version 3.2
;; SEQ ID NO 2
;; LENGTH: 145
;; TYPE: PRT
;; ORGANISM: Homo sapiens
US-09-528-436B-2
Query Match 25.1%; Score 165.5; DB 4; Length 145;
Best Local Similarity 32.5%; Pred. No. 1e-10;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;
Qy 9 TKICVGPDRPTNSPELEETLTTITIKLNAENATYFKIDNVKARQVAVGKYYFID 68
Db 32 SRVKGPFPTIKTNDPGVLQAARYSVEKFNNTNDMFLKESRITRALVOIVKGLKYMLE 91
Qy 69 FVARETTCKESNEELTESCS--TKLQSLDCNAEVVVPWEKKI-YPTVNC 118
Db 92 VEIGRTCKCKQHRL-LDCCDFQTNHILKQTLSCYSEVVVPWLQHPVPVLR 144
RESULT 11
US-08-676-242-11
;; Sequence 11, Application US/08676242C
;; Patent No. 6143719
;; GENERAL INFORMATION:
;; APPLICANT: The Regents of the University of Michigan
;; APPLICANT: Schmaier, Alvin H.
;; APPLICANT: Hasan, Ahmed A.K.
;; TITLE OF INVENTION: Bradykinin Analogs As Selective Thrombin Inhibitors
;; FILE REFERENCE: 8820-2 US
;; CURRENT APPLICATION NUMBER: US/08/676,242C
;; PRIOR FILING DATE: 2000-07-16
;; EARLIER APPLICATION NUMBER: 60/000,096
;; EARLIER FILING DATE: 1995-06-09
;; PRIOR FILING DATE: 1996-06-07
;; NUMBER OF SEQ ID NOS: 24
;; SOFTWARE: Patent in Ver. 2.1
;; SEQ ID NO 11
;; LENGTH: 27
;; TYPE: PRT
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: Description of Artificial Sequence: Bradykinin
US-08-676-242-11
Query Match 23.5%; Score 155; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 1.6e-10;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 97 LDCNAEVVVPWEKKIYPTVNCQPLGM 123
Db 1 LDCNAEVVVPWEKKIYPTVNCQPLGM 27
RESULT 12
US-08-676-242-22
;; Sequence 22, Application US/08676242C
;; Patent No. 6143719
;; GENERAL INFORMATION:
;; APPLICANT: The Regents of the University of Michigan
;; APPLICANT: Schmaier, Alvin H.
;; APPLICANT: Hasan, Ahmed A.K.
;; TITLE OF INVENTION: Bradykinin Analogs As Selective Thrombin Inhibitors
;; FILE REFERENCE: 8820-2 US
;; CURRENT APPLICATION NUMBER: US/08/676,242C

; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 14
; LENGTH: 121
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-14

Query Match 21.0%; Score 138.5; DB 4; Length 121;
Best Local Similarity 31.5%; Pred. No. 7.7e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
QY 8 PTKICVCCPRDIPITNSPELEETLTHITKLNANNATFYFKIDNVKARVQVWAGKKYFI 67
DB 2 PQERMVGEIRDLSPPDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSQLVAGIKYFL 61
QY 68 DFVARETTCSKE---SNEELTESCETKKGQ--SLDCNAEVVVPWE 109
DB 62 TMEMGSTDCKTRVTGHDVLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 108

RESULT 15

US-09-775-932-12
; Sequence 12, Application US/09775932
; Patent No. 6534477
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 12
; LENGTH: 128
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-12

Query Match 21.0%; Score 138.5; DB 4; Length 128;
Best Local Similarity 31.5%; Pred. No. 8.3e-08;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
QY 8 PTKICVCCPRDIPITNSPELEETLTHITKLNANNATFYFKIDNVKARVQVWAGKKYFI 67
DB 9 PQERMVGEIRDLSPPDPQVQKAAQAAVASYNMGNSIYFRDTHIIKAQSQLVAGIKYFL 68
QY 68 DFVARETTCSKE---SNEELTESCETKKGQ--SLDCNAEVVVPWE 109
DB 69 TMEMGSTDCKTRVTGHDVLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 115

Search completed: September 24, 2004, 14:11:36
Job time : 27.268 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:02:27 ; Search time 49.692 Seconds
(without alignments)
699.375 Million cell updates/sec

Title: US-10-661-784-1
Perfect score: 660
Sequence: 1 GKDFVQPTKICVGCPRDIP.....VVPWEKKIYPTVNCQLGM 123

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_29Jan04:*

- 1: Geneseq1980s:*
- 2: Geneseq1990s:*
- 3: Geneseq2000s:*
- 4: Geneseq2001s:*
- 5: Geneseq2002s:*
- 6: Geneseq2003as:*
- 7: Geneseq2003bs:*
- 8: Geneseq2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	660	100.0	123	3	AA95426	Human hig
2	660	100.0	304	6	ABP70801	Human ext
3	660	100.0	322	6	ABP70799	Human ext
4	660	100.0	329	6	ABU92044	Human pro
5	660	100.0	358	6	ABP70800	Human ext
6	660	100.0	390	6	ABU99149	Novel hum
7	660	100.0	398	6	ABU99143	Novel hum
8	660	100.0	427	8	ADE76864	Human pro
9	660	100.0	615	6	ABU99144	Novel hum
10	660	100.0	626	5	ABP78707	Human hig
11	660	100.0	644	4	ABG21101	Novel hum
12	660	100.0	644	5	ABP78710	Human hig
13	660	100.0	644	6	ABU99150	Novel hum
14	660	100.0	644	6	ABU99145	Novel hum
15	628	95.2	122	3	AA95426	Human hig
16	594	90.0	117	2	AA933350	Domaine 3
17	589.5	89.3	435	4	ABG21105	Novel hum
18	477	72.3	436	1	AA940257	Bradykini
19	453	68.6	357	6	ABR41202	Human DIT
20	450	68.2	434	1	AA940633	Human DIT
21	409	62.0	235	5	ABG60077	Human DIT
22	320.5	48.6	248	4	ABG21102	Novel hum
23	316	47.9	369	4	ABG21099	Novel hum
24	190	28.8	305	4	ABG21100	Novel hum
25	175	26.5	167	2	AA938907	Mouse IMC

26	169	25.6	178	2	AAW69734	Human cys
27	166	25.2	32	3	AA95418	Anti-angi
28	165.5	25.1	126	3	AB37445	Human cys
29	165.5	25.1	145	2	AAW32323	Mature hu
30	165.5	25.1	145	2	AAW31502	Human cys
31	165.5	25.1	145	2	AAW31502	Human cys
32	165.5	25.1	145	4	AAE02410	Human cys
33	165.5	25.1	145	4	AAE04439	Human cys
34	165.5	25.1	145	7	ADD14098	Human src
35	165.5	25.1	167	7	AAV02287	Secreted
36	165.5	25.1	167	7	ADA45154	Human pol
37	161	24.4	32	3	AA95408	Anti-angi
38	157	23.8	122	3	AB37446	Human kin
39	156.5	23.7	167	2	AAW98910	Mouse IMC
40	155	23.5	27	2	AAW54335	Bradykini
41	153	23.2	27	3	AA95425	Anti-angi
42	145	22.0	64	2	AAW54341	Bradykini
43	145	22.0	64	2	AAW77418	Kininogen
44	144.5	21.9	121	3	AAV81200	Human mut
45	144.5	21.9	128	3	AAV81189	Human mut

ALIGNMENTS

RESULT 1
AA95426
ID AA95426 standard; peptide; 123 AA.
XX
AC AA95426;
XX
DT 25-SEP-2000 (first entry)
XX
DE Human high mol.wt. kininogen domain 3.
XX
KW Anti-angiogenic; angiogenesis; inhibitor; kininogen; homologue;
KW endothelial cell proliferation; apoptosis; cancer; ocular disorder;
KW rheumatoid arthritis; cytostatic; antiarthritic; antirheumatic; therapy;
KW human; D3 peptide.
XX
OS Homo sapiens.
XX
PN WO200035407-A2.
XX
PD 22-JUN-2000.
XX
PF 02-DEC-1999; 99WO-US028465.
XX
PR 16-DEC-1998; 98US-0112427P.
XX
PA (UTEM) UNIV TEMPLE.
XX
PI (MCCR/) MCCR R K.
XX
DR Mcrae RK;
XX
WPI; 2000-442247/38.
XX
PT Composition for inhibiting angiogenesis and endothelial cell
PT proliferation, inducing endothelial cell apoptosis and treating cancer, 3
PT rheumatoid arthritis, and ocular disorders comprises a kininogen domain 3
XX
PS Disclosure; Page 4; 44pp; English.
XX
CC The present sequence is that of domain 3 of human high mol.wt. kininogen
CC (HK). The invention provides peptides (see AA95405-24) that are
CC analogues of certain sites in the HK domain 3, specifically Asn275-
CC Lys282, Cys246-Cys249, Leu331-Tyr338 and Tyr299-Ser314. The peptides, in
CC which native Cys residues may be replaced by Ala residues, inhibit
CC endothelial cell proliferation and may also induce endothelial cell
CC apoptosis. Compositions including the peptides are used in claimed
CC methods for inhibiting angiogenesis, inhibiting endothelial cell
CC proliferation, and inducing endothelial cell apoptosis. Cancer.

CC rheumatoid arthritis, and ocular disorders characterized by undesired
 CC vascularization of the retina are treated
 XX
 SQ Sequence 123 AA;
 Query Match 100.0%; Score 660; DB 3; Length 123;
 Best Local Similarity 100.0%; Pred. No. 2.3e-66;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLNAENNAFFYFKIDNVKKARQVQV 60
 DB 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLNAENNAFFYFKIDNVKKARQVQV 60
 QY 61 AGKXYFIDFVARETTCSKESNEELTSCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 120
 DB 61 AGKXYFIDFVARETTCSKESNEELTSCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 120
 QY 121 LGM 123
 DB 121 LGM 123
 RESULT 2
 ABP70801
 ID ABP70801 standard; protein; 304 AA.
 XX
 AC ABP70801;
 XX
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-28.
 XX
 DE Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.
 XX
 PA (INCY-) INCYTE GENOMICS INC.
 XX
 PI Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebajadian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Ramkumar J;
 XX
 DR WPI; 2003-278643/27.
 DR N-PSDB; ACC42388.
 XX
 XX New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 disorders or cancer.
 PT
 XX
 PS Claim 1; Page 207; 224pp; English.
 XX
 XX The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP70774-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition

CC for diagnosing or treating a disease or condition associated with.
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer
 XX
 SQ Sequence 304 AA;
 Query Match 100.0%; Score 660; DB 6; Length 304;
 Best Local Similarity 100.0%; Pred. No. 8e-66;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLNAENNAFFYFKIDNVKKARQVQV 60
 DB 130 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLNAENNAFFYFKIDNVKKARQVQV 189
 QY 61 AGKXYFIDFVARETTCSKESNEELTSCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 120
 DB 190 AGKXYFIDFVARETTCSKESNEELTSCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 249
 QY 121 LGM 123
 DB 250 LGM 252
 RESULT 3
 ABP70799
 ID ABP70799 standard; protein; 322 AA.
 XX
 AC ABP70799;
 XX
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-26.
 XX
 DE Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.
 XX
 PA (INCY-) INCYTE GENOMICS INC.
 XX
 PI Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebajadian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Ramkumar J;
 XX
 DR WPI; 2003-278643/27.
 DR N-PSDB; ACC42386.
 XX
 XX New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 disorders or cancer.
 PT
 XX
 PS Claim 1; Page 205-206; 224pp; English.
 XX
 XX The present invention relates to novel human extracellular messenger

CC proteins (EXMES-1 to-28; ABP70774-ABP70801) and their coding sequences
CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
CC for diagnosing or treating a disease or condition associated with
CC decreased expression or overexpression of functional EXMES e.g.
CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
CC cancer
XX
SQ Sequence 322 AA;
Query Match 100.0%; Score 660; DB 6; Length 322;
Best Local Similarity 100.0%; Pred. No. 8.7e-66; Indels 0; Gaps 0;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITIKLNAENNAFFYKIDNVKKARQVW 60
DB 148 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITIKLNAENNAFFYKIDNVKKARQVW 207
QY 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTVNCQP 120
DB 208 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTVNCQP 267
QY 121 LGM 123
DB 268 LGM 270
RESULT 4
ID ABU92044 standard; protein; 329 AA.
XX
AC ABU92044;
XX
DT 15-JUL-2003 (first entry)
XX
DE Human protein modification and maintenance molecule-24 (PMM-24).
KW Human; protein modification and maintenance molecule; PMM; cancer;
KW cell proliferation disorder; atherosclerosis; neurological disorder;
KW epilepsy; Huntington's disease; stroke; immune disorder; allergy;
KW inflammatory disorder; AIDS; developmental disorder; hypothyroidism;
KW Cushing's syndrome; gastrointestinal disorder; epithelial disorder;
KW infection; cytostatic; antiarteriosclerotic; anticonvulsant; nootropic;
KW neuroprotective; cerebroprotective; anti-HIV; antiallergic; vulnary;
KW antiinflammatory; thyromimetic.
XX
OS Homo sapiens.
XX
XX WO2003031939-A2.
XX
PD 17-APR-2003.
XX
XX 11-OCT-2002; 2002WO-US032850.
XX
PR 12-OCT-2001; 2001US-0329689P.
PR 25-OCT-2001; 2001US-0335703P.
PR 09-NOV-2001; 2001US-0346887P.
PR 28-NOV-2001; 2001US-0334145P.
PR 06-DEC-2001; 2001US-0337451P.
PR 14-DEC-2001; 2001US-0340584P.
XX
XX (INCY-) INCYTE GENOMICS INC.
XX
PI Ramkumar J, Gorvad AE, Baughn MR, Emerling BM, Yang J, Lee SY;
PI Tran UK, Becha SD, Duggan BM, Lee EA, Griffin JA, Li JX;
PI Sprague WW, Rafalia AJA, Chawla NK, Lehr-Mason PM, Kable AE, Yue H;
PI Marquis JP, Yao MG, Richardson TW, Tang TY, Jin P, Chien D;
PI Bhatia U, Burrill JD, Lee S, Blake JJ, Ho A, Zheng W;
XX
XX WPI; 2003-430274/40.
DR N-PSDB; ACA92439.
XX
PT New human protein modification and maintenance molecules (PMM), useful
PT for diagnosing, treating and preventing diseases or conditions associated

PT with the aberrant PMM expression e.g. cancer, atherosclerosis, or
PT infections.
XX
PS Claim 1; Page 264-265; 311pp; English.
XX
CC The present invention relates to the isolation of human protein
CC modification and maintenance molecules (PMM), and the polynucleotide
CC sequences encoding them. A total of 40 PMM polypeptides (designated PMM
CC -1 to PMM-40) are disclosed. The sequences of the invention are useful
CC for diagnosing a condition or disease associated with the expression of
CC PMM in a subject, preparing a polyclonal or monoclonal antibody, and
CC generating an expression profile of a sample containing the
CC polynucleotides. The diseases or conditions associated with decreased
CC expression or overexpression of PMM are cell proliferation disorders
CC (e.g. cancer, atherosclerosis), neurological disorders (e.g. epilepsy,
CC Huntington's disease, stroke), immune/inflammatory disorders, (e.g. AIDS,
CC allergies), developmental disorders (e.g. hypothyroidism, Cushing's
CC syndrome), gastrointestinal disorders (e.g. epithelial disorders, and infections. The
CC PMM polypeptides or their fragments are useful in screening compounds
CC for effectiveness as agonists or antagonists of the polypeptides, or in
CC altering the expression of the target polynucleotide and compounds that
CC specifically bind to, or modulate the activity of the polypeptide.
CC ABU92021-ABU92060 represent the human PMM polypeptides of the invention
XX
SQ Sequence 329 AA;
Query Match 100.0%; Score 660; DB 6; Length 329;
Best Local Similarity 100.0%; Pred. No. 9e-66;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITIKLNAENNAFFYKIDNVKKARQVW 60
DB 155 GKDFVQPTKICVGCPRDIPITNSPELEETLTHITIKLNAENNAFFYKIDNVKKARQVW 214
QY 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTVNCQP 120
DB 215 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEYVVPWEKKIYPTVNCQP 274
QY 121 LGM 123
DB 275 LGM 277
RESULT 5
ID ABP70800 standard; protein; 358 AA.
XX
AC ABP70800;
XX
DT 26-AUG-2003 (first entry)
XX
DE Human extracellular messenger, EXMES-27.
XX
KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
KW endocrine disorder; cancer.
XX
OS Homo sapiens.
XX
XX WO2003018612-A2.
XX
XX 06-MAR-2003.
XX
XX 22-AUG-2002; 2002WO-US027213.
XX
PR 24-AUG-2001; 2001US-0314811P.
PR 14-DEC-2001; 2001US-0340584P.
PR 18-JAN-2002; 2002US-0350595P.
PR 11-MAR-2002; 2002US-0363432P.
PR 15-MAR-2002; 2002US-0364607P.
PR 05-APR-2002; 2002US-0370761P.
PR 24-JUN-2002; 2002US-0391378P.
XX

PA (INCY-) INCYTE GENOMICS INC.
 XX Duggan BM, Lee S, Raughn MR, Hafalia AJA, Walia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebajadian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Ramkumar J;
 XX WPI: 2003-278643/27.
 DR N-PSDB; ACC42387.
 XX
 PT New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 206; 224pp; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP70774-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
 CC for diagnosing or treating a disease or condition associated with
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer
 XX
 SQ Sequence 358 AA;
 Query Match 100.0%; Score 660; DB 6; Length 358;
 Best Local Similarity 100.0%; Pred. No. 1e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPPTKICVGCPRDIPPTNSPELEETLTHITIKLNAENNAFFKIDNVKKARVQV 60
 DB 184 GKDFVQPPTKICVGCPRDIPPTNSPELEETLTHITIKLNAENNAFFKIDNVKKARVQV 243
 QY 61 AGKKYFIDFVARETTCESKSNELTSCCTKLGSLDCNAEYVVPWEKKIYPTVNCQP 120
 DB 244 AGKKYFIDFVARETTCESKSNELTSCCTKLGSLDCNAEYVVPWEKKIYPTVNCQP 303
 QY 121 LGM 123
 DB 304 LGM 306
 RESULT 6
 ABU99149
 AC ABU99149 standard; protein; 390 AA.
 XX ABU99149;
 DT 01-AUG-2003 (first entry)
 XX
 DE Novel human GPCR related protein NOV129.
 XX
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiac; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiaesthmic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX
 OS Homo sapiens.
 XX
 XX WO200299116-A2.
 FN
 XX
 PD 12-DEC-2002.
 XX

PF 04-JUN-2002; 2002WO-US017428.
 XX 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295661P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 06-JUN-2001; 2001US-0296418P.
 PR 14-JUN-2001; 2001US-0298285P.
 PR 15-JUN-2001; 2001US-0298556P.
 PR 21-JUN-2001; 2001US-0299949P.
 PR 26-JUN-2001; 2001US-0300883P.
 PR 28-JUN-2001; 2001US-0301550P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 27-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-0322293P.
 PR 17-SEP-2001; 2001US-0322706P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR 03-JUN-2002; 2002US-00363676.
 XX (CURA-) CURAGEN CORP.
 PA Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;
 PI Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalte T, Kekuda R, Li L;
 PI Macdougall JR, Malyankar UM, Millet I, Padigaru M, Paturajan M;
 PI Pena CE, Rasteili L, Shinkets RA, Stone DU, Spytek KA, Vernet CAM;
 PI Voss EZ, Zerhusen BD;
 XX WPI: 2003-140627/13.
 DR N-PSDB; AC03653.
 XX
 PT New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 PS Claim 1; Page 147; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 SQ Sequence 390 AA;
 Query Match 100.0%; Score 660; DB 6; Length 390;
 Best Local Similarity 100.0%; Pred. No. 1.1e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPPTKICVGCPRDIPPTNSPELEETLTHITIKLNAENNAFFKIDNVKKARVQV 60
 DB 216 GKDFVQPPTKICVGCPRDIPPTNSPELEETLTHITIKLNAENNAFFKIDNVKKARVQV 275

QY 61 AGKYFIDFVARETTCKESNEELTESCETKLGSLDCNAEYVYVWPWEKKIYPTVNCQP 120
 DB 276 AGKYFIDFVARETTCKESNEELTESCETKLGSLDCNAEYVYVWPWEKKIYPTVNCQP 335
 QY 121 LGM 123
 DB 336 LGM 338
 RESULT 7
 ABUS9143
 ID ABUS9143 standard; protein; 398 AA.
 AC ABUS9143;
 XX
 DT 01-AUG-2003 (first entry)
 XX
 DE Novel human GPCR related protein NOV12a.
 XX
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiant; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOXV-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX
 OS Homo sapiens.
 XX
 WO200299116-A2.
 XX
 PN 12-DEC-2002.
 XX
 PD 04-JUN-2002; 2002WO-US017428.
 XX
 PF 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295661P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 06-JUN-2001; 2001US-0296418P.
 PR 14-JUN-2001; 2001US-0298285P.
 PR 15-JUN-2001; 2001US-0298558P.
 PR 21-JUN-2001; 2001US-0299949P.
 PR 26-JUN-2001; 2001US-0300883P.
 PR 28-JUN-2001; 2001US-0301550P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 27-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-0322293P.
 PR 17-SEP-2001; 2001US-0322708P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR 03-JUN-2002; 2002US-00363676.
 XX
 PA (CURA-) CURAGEN CORP.
 XX
 PI Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;
 PI Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalte T, Kekuda R, Li L;
 PI MacDougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
 PI Pena CE, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CM;
 PI Voss EZ, Zehrhusen BD;
 XX
 WPI; 2003-140627/13.
 DR N-PSDB; ACD03647.
 DR
 XX
 XX New NOXV polypeptides and nucleic acids, useful for preventing or
 PT treating NOXV-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or

PT pharmacogenomics.
 XX
 PS Claim 1; Page 143; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOXV polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOXV-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOXV substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 SQ Sequence 398 AA;
 Query Match 100.0%; Score 660; DB 6; Length 398;
 Best Local Similarity 100.0%; Pred. No. 1.2e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLNANNATFYFKIDNVKKARQVYV 60
 DB 224 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLNANNATFYFKIDNVKKARQVYV 283
 QY 61 AGKYFIDFVARETTCKESNEELTESCETKLGSLDCNAEYVYVWPWEKKIYPTVNCQP 120
 DB 284 AGKYFIDFVARETTCKESNEELTESCETKLGSLDCNAEYVYVWPWEKKIYPTVNCQP 343
 QY 121 LGM 123
 DB 344 LGM 346
 RESULT 8
 ADE76864
 ID ADE76864 standard; protein; 427 AA.
 XX
 AC ADE76864;
 XX
 DT 29-JAN-2004 (first entry)
 XX
 DE Human protein expressed in a liver disorder #9.
 XX
 KW human; liver disorder; hyperlipidaemia; hypertension; type II diabetes;
 KW tumour; liver; inflammatory disorder; immune response disorder;
 KW high-throughput screening; differential gene expression; gene therapy.
 XX
 OS Homo sapiens.
 XX
 FN US2003108871-A1.
 PD 12-JUN-2003.
 XX
 FF 30-JUL-2001; 2001US-00919039.
 XX
 XX 28-JUL-2000; 2000US-0222113P.
 XX
 PA (KASE/) KASER M R.
 XX
 PI Kaser MR;

XX WPI; 2004-031227/03.
 DR N-PSDB; ADE76863.
 XX
 PT Composition comprising several cDNAs that are differentially expressed in
 PT treated human C3A liver cell cultures, useful for treating liver
 PT disorders.
 XX
 PS Claim 1; SEQ ID NO 29; 41pp; English.
 XX
 CC The invention relates to a composition comprising several cDNAs that are
 CC differentially expressed in a liver disorder. The composition is useful
 CC for treating liver disorder such as hyperlipidaemia, hypertension, type
 CC II diabetes, tumours of the liver and disorders of the inflammatory and
 CC immune response. The composition is useful for a high-throughput method
 CC of screening several molecules or compounds to identify a ligand which
 CC specifically binds a cDNA. A protein encoded by the cDNA is useful for a
 CC high-throughput method for using a protein to screen several molecules or
 CC compounds to identify at least one ligand which specifically binds the
 CC protein which involves combining the protein encoded by the cDNA with
 CC several of molecules or compounds under conditions to allow specific
 CC binding, and detecting specific binding between the protein and a
 CC molecule or compound, therefore identifying a ligand which specifically
 CC binds the protein. The composition is useful for detecting and
 CC quantifying differential gene expression, can be used in gene therapy, to
 CC formulate prognosis and to design a treatment regimen and to monitor the
 CC efficacy of treatment. The present sequence represents the amino acid
 CC sequence of a protein encoded by a cDNA differentially expressed in a
 CC liver disorder.
 XX
 XX Sequence 427 AA;
 Query Match 100.0%; Score 660; DB 8; Length 427;
 Best Local Similarity 100.0%; Pred. NO. 1.3e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPPKICVGCPRDPTNSPELEETLTHITKLAENNATVFKIDNVKKARVQV 60
 DB 253 GKDFVQPPKICVGCPRDPTNSPELEETLTHITKLAENNATVFKIDNVKKARVQV 312
 QY 61 AGKXIFDFARETTCKSNBELTESCTKLGSLDCNAEYVVPWEKKIYPTVNCQ 120
 DB 313 AGKXIFDFARETTCKSNBELTESCTKLGSLDCNAEYVVPWEKKIYPTVNCQ 372
 QY 121 LGM 123
 DB 373 LGM 375
 RESULT 9
 ID ABU99144
 AC ABU99144 standard; protein; 615 AA.
 XX
 XX ABU99144;
 XX
 DT 01-AUG-2003 (first entry)
 DE Novel human GPCR related protein NOV12b.
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiac; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX Homo sapiens.
 OS

PN W0200299116-A2.
 XX
 PD 12-DEC-2002.
 XX
 PF 04-JUN-2002; 2002WO-US017428.
 PR 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295651P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 06-JUN-2001; 2001US-0296418P.
 PR 14-JUN-2001; 2001US-0298285P.
 PR 15-JUN-2001; 2001US-0298556P.
 PR 21-JUN-2001; 2001US-0299949P.
 PR 26-JUN-2001; 2001US-0300893P.
 PR 28-JUN-2001; 2001US-0301550P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 27-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-0322293P.
 PR 17-SEP-2001; 2001US-0322706P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR 03-JUN-2002; 2002US-00363676.
 XX (CURA-) CURAGEN CORP.
 PA
 XX Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR,
 PI Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalte T, Kekuda R, Li L;
 PI Macdougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
 PI Pena CE, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CM;
 PI Voss EZ, Zernhosen BD;
 XX WPI; 2003-140627/13.
 DR N-PSDB; ACD03648.
 XX
 XX New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 PS Claim 1; Page 144; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 XX Sequence 615 AA;
 Query Match 100.0%; Score 660; DB 6; Length 615;
 Best Local Similarity 100.0%; Pred. NO. 2.1e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGCPRDIPNTSPSELEETLTHITITKLNNAENNAFFYFKIDNVKKARQVQV 60
Db 224 GKDFVQPTKICVGCPRDIPNTSPSELEETLTHITITKLNNAENNAFFYFKIDNVKKARQVQV 283
QY 61 AGKYFIDFVARETTCSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTVNCOP 120
Db 284 AGKYFIDFVARETTCSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTVNCOP 343
QY 121 LGM 123
Db 344 LGM 346
RESULT 10
ABB78707
ID ABB78707 standard; protein; 626 AA.
XX
AC ABB78707;
XX
DT 18-JUL-2002 (first entry)
XX
DE Human high molecular weight kininogen (HK) mature protein SEQ ID NO:1.
XX
KW Human; kininogen; high molecular weight kininogen; HK; D5 domain;
KW D5 receptor; angiogenesis; endothelial cell; cytostatic; antitumour;
KW antiatherosclerotic; vasotropic; vulnery; tranquilliser; thrombolytic;
KW ophthalmological; gynaecological; antiulcer; antidiabetic; antiarthritic;
KW antiangiogenic; antiapoptotic; endocrine; apoptosis; gene therapy.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT Domain 384..508
FT /label= D5_domain
XX
FN WO200214369-A2.
XX
PD 21-FEB-2002;
XX
PF 24-JUL-2001; 2001WO-US023185.
XX
PR 24-JUL-2000; 2000US-0220194P.
XX
FA (ATTE-) ATTENUON LLC.
XX
PI Mazar AP, Juarez JC;
XX
XX WPI; 2002-393611/42.
XX
PT Novel human kininogen D5 domain polypeptides useful for treating
PT conditions associated with endothelial cell migration, proliferation,
PT invasion or angiogenesis, e.g. arthritis, macular degeneration, benign
PT hyperplasia.
XX
PS Disclosure; Page 13; 84pp; English.
XX
CC The present invention describes an isolated polypeptide (I) that
CC corresponds to the D5 domain of human kininogen, or biologically active
CC peptide fragment, homologue or functional derivative, and which: (a)
CC inhibits angiogenesis; (b) binds to the D5 binding site on endothelial
CC cells (EC); (c) activates signalling pathways leading to the introduction
CC of apoptosis in EC; and/or (d) inhibits the signalling pathway required
CC for maintenance of EC viability. (I) has cytostatic, antitumour,
CC antiatherosclerotic, vasotropic, vulnery, tranquilliser, thrombolytic,
CC ophthalmological, gynaecological, antiulcer, antidiabetic, antiarthritic,
CC antiangiogenic, antiapoptotic and endocrine activities. An antibody (IX)
CC specific for an epitope of (I) is useful for inhibiting tumour growth or
CC angiogenesis in a subject. (II), a D5 fusion polypeptide (II) or a dimeric
CC or trimeric fusion polypeptide (III) can be used for inhibiting EC
CC migration, proliferation, invasion, or angiogenesis, or for inducing EC
CC apoptosis. An angiogenic EC-targeting pharmaceutical composition (X)
CC comprising (I), (II), or (III), can be used for treating a subject having
CC a disease or condition associated with undesired EC migration,

CC proliferation, invasion or angiogenesis. (I), (II), or (III) can be used
CC for isolating a D5 domain binding molecule from a complex mixture and for
CC isolating or enriching cells expressing D5 domain binding sites from a
CC cell mixture. The present sequence represents the mature human high
CC molecular weight kininogen (HK) protein, which is given in the
CC exemplification of the present invention
XX
SQ Sequence 626 AA;

Query Match 100.0%; Score 660; DB 5; Length 626;
Best Local Similarity 100.0%; Pred. No. 2.2e-65;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGCPRDIPNTSPSELEETLTHITITKLNNAENNAFFYFKIDNVKKARQVQV 60
Db 235 GKDFVQPTKICVGCPRDIPNTSPSELEETLTHITITKLNNAENNAFFYFKIDNVKKARQVQV 294
QY 61 AGKYFIDFVARETTCSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTVNCOP 120
Db 295 AGKYFIDFVARETTCSKESNEELTESCETKKLGSLDCNAEVVVPWEKKIYPTVNCOP 354
QY 121 LGM 123
Db 355 LGM 357

RESULT 11

ID ABB21101 standard; protein; 644 AA.
XX

AC ABB21101;

DT 18-FEB-2002 (first entry)

DE Novel human diagnostic protein #21092.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.

XX Homo sapiens.

XX WO200175067-A2.

PN 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US008631.

PR 31-MAR-2000; 2000US-00540217.

PR 23-AUG-2000; 2000US-00649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

XX N-PSDB; AAS95288.

XX New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity.

XX Claim 20; SEQ ID NO 51460; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and polypeptide (II)
CC sequences. (I) is useful as hybridisation probes, polymerase chain
CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
CC and in recombinant production of (II). The polynucleotides are also used
CC in diagnostics as expressed sequence tags for identifying expressed
CC genes. (I) is useful in gene therapy techniques to restore normal
CC activity of (II) or to treat disease states involving (II). (II) is
CC useful for generating antibodies against it, detecting or quantitating a
CC polypeptide in tissue, as molecular weight markers and as a food

CC supplement. (II) and its binding partners are useful in medical imaging
 CC of sites expressing (II). (I) and (II) are useful for treating disorders
 CC involving aberrant protein expression or biological activity. The
 CC polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. ABG00010-ABG30377 represent novel human diagnostic
 CC amino acid sequences of the invention. Note: The sequence data for this
 CC patent did not appear in the printed specification, but was obtained in
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 XX Sequence 644 AA;

Query Match 100.0%; Score 660; DB 4; Length 644;
 Best Local Similarity 100.0%; Pred. No. 2.3e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPPKICVGCPRDPTNSPELEETLTHITKLNAENNAFFYFKIDNVKARQVW 60
 Db 253 GKDFVQPPKICVGCPRDPTNSPELEETLTHITKLNAENNAFFYFKIDNVKARQVW 312
 QY 61 AGKKYFIDFVARETTCKESNEELTESCTKKLGSLDCNAEYVVPWEKKIYPTVNCOP 120
 Db 313 AGKKYFIDFVARETTCKESNEELTESCTKKLGSLDCNAEYVVPWEKKIYPTVNCOP 372
 QY 121 LGM 123
 Db 373 LGM 375

RESULT 12
 ID ABB78710
 XX ABB78710 standard; protein; 644 AA.
 AC ABB78710;
 XX
 DT 18-JUL-2002 (first entry)
 XX Human high molecular weight kininogen (HK) protein.
 DE Human; kininogen; high molecular weight kininogen; HK; D5 domain;
 KW D5 receptor; angiogenesis; endothelial cell; cytostatic; antitumour;
 KW antiatherosclerotic; vasotropic; vulnary; tranquilliser; thrombolytic;
 KW ophthalmological; gynaecological; antiulcer; antidiabetic; antiarthritic;
 KW antiangiogenic; antiapoptotic; endocrine; apoptosis; gene therapy.
 XX Homo sapiens.

Key	Location/Qualifiers
Peptide	1..18
Protein	/label= signal
	13..644
	/label= mature_human_high_molecular_weight_kininogen
Disulfide-bond	28..614
Disulfide-bond	83..94
Disulfide-bond	107..126
Disulfide-bond	142..145
Disulfide-bond	206..218
Disulfide-bond	229..248
Disulfide-bond	264..267
Disulfide-bond	328..340
Disulfide-bond	351..370
Domain	402..526
	/label= D5_domain

WO200214369-A2.

21-FEB-2002.

24-JUL-2001; 2001WO-US023185.

PR 24-JUL-2000; 2000US-0220194P.
 XX (ATTE-) ATTENUON LLC.
 PA Mazar AP, Juarez JC;
 FI WPI; 2002-393611/42.
 XX Novel human kininogen D5 domain polypeptides useful for treating
 PT conditions associated with endothelial cell migration, proliferation,
 PT invasion or angiogenesis, e.g. arthritis, macular degeneration, benign
 PT hyperplasia.
 XX Disclosure; Fig 1B-E; 84pp; English.
 PS
 XX The present invention describes an isolated polypeptide (I) that
 CC corresponds to the D5 domain of human kininogen, or biologically active
 CC peptide fragment, homologue or functional derivative, and which (a)
 CC inhibits angiogenesis; (b) binds to the D5 binding site on endothelial
 CC cells (EC); (c) activates signalling pathways leading to the introduction
 CC of apoptosis in EC; and/or (d) inhibits the signalling pathway required
 CC for maintenance of EC viability. (I) has cytostatic, antitumour,
 CC antiatherosclerotic, vasotropic, vulnary, tranquilliser, thrombolytic,
 CC ophthalmological, gynaecological, antiulcer, antidiabetic, antiarthritic,
 CC antiangiogenic, antiapoptotic and endocrine activities. An antibody (IX)
 CC specific for an epitope of (I) is useful for inhibiting tumour growth or
 CC angiogenesis in a subject. (I), a D5 fusion polypeptide (II) or a dimeric
 CC or trimeric fusion polypeptide (III) can be used for inhibiting EC
 CC migration, proliferation, invasion, or angiogenesis, or for inducing EC
 CC apoptosis. An angiogenic EC-targeting pharmaceutical composition (X)
 CC comprising (I), (II), or (III), can be used for treating a subject having
 CC a disease or condition associated with undesired EC migration,
 CC proliferation, invasion or angiogenesis. (I), (II), or (III) can be used
 CC for isolating a D5 domain binding molecule from a complex mixture and for
 CC isolating or enriching cells expressing D5 domain binding sites from a
 CC cell mixture. The present sequence represents the human high molecular
 CC weight kininogen (HK) protein, which is given in the exemplification of
 CC the present invention
 XX
 SQ Sequence 644 AA;

Query Match 100.0%; Score 660; DB 5; Length 644;
 Best Local Similarity 100.0%; Pred. No. 2.3e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPPKICVGCPRDPTNSPELEETLTHITKLNAENNAFFYFKIDNVKARQVW 60
 Db 253 GKDFVQPPKICVGCPRDPTNSPELEETLTHITKLNAENNAFFYFKIDNVKARQVW 312
 QY 61 AGKKYFIDFVARETTCKESNEELTESCTKKLGSLDCNAEYVVPWEKKIYPTVNCOP 120
 Db 313 AGKKYFIDFVARETTCKESNEELTESCTKKLGSLDCNAEYVVPWEKKIYPTVNCOP 372
 QY 121 LGM 123
 Db 373 LGM 375

RESULT 13

ABU99150
 ID ABU99150 standard; protein; 644 AA.

XX ABU99150;

XX 01-AUG-2003 (first entry)

XX Novel human GPCR related protein NOV12h.

DE Human; G-protein coupled receptor related protein; GPCR related protein;
 XX NOV; cytostatic; cardiant; antiatherosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;

KW diabetes; immune disorder; AIDS; obesity; asthma;
KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
KW infection; multiple sclerosis; cancer-associated cachexia;
KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
XX Homo sapiens.
XX WO200299116-A2.
XX PD 12-DEC-2002.
XX 04-JUN-2002; 2002WO-US017428.
XX 04-JUN-2001; 2001US-0295607P.
PR 04-JUN-2001; 2001US-0295661P.
PR 06-JUN-2001; 2001US-0296404P.
PR 06-JUN-2001; 2001US-0296418P.
PR 14-JUN-2001; 2001US-0298285P.
PR 15-JUN-2001; 2001US-0298566P.
PR 21-JUN-2001; 2001US-0299949P.
PR 26-JUN-2001; 2001US-0300883P.
PR 28-JUN-2001; 2001US-0301550P.
PR 13-AUG-2001; 2001US-0311972P.
PR 27-AUG-2001; 2001US-0315071P.
PR 29-AUG-2001; 2001US-0315660P.
PR 14-SEP-2001; 2001US-0322293P.
PR 17-SEP-2001; 2001US-0322706P.
PR 14-DEC-2001; 2001US-0341186P.
PR 17-SEP-2001; 2001US-0322293P.
PR 14-DEC-2001; 2001US-0341186P.
PR 28-FEB-2002; 2002US-0361189P.
PR 12-MAR-2002; 2002US-0363673P.
PR 12-MAR-2002; 2002US-0363676P.
PR 03-JUN-2002; 2002US-00363676.
XX (CURA-) CURAGEN CORP.
XX Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR, Li L;
PI Gangolli EA, Gerlach VL, Gorman L, Hjalit T, Kekuda R, Li L;
PI Macdougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
PI Pena CBA, Rastelli L, Shimkets RA, Stone DJ, Spytek KA, Vernet CAM;
PI Voss EZ, Zerhusen BD;
XX WPI; 2003-140627/13.
DR N-PSDB; ACD03654.
XX New NOVX polypeptides and nucleic acids, useful for preventing or
PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
PT pharmacogenomics.
XX Claim 1; Page 148; 332pp; English.
XX The invention describes an isolated polypeptide (I) comprising any of 27
CC 118-961 residue amino acid sequences, given in the specification, a
CC mature form of them, a sequence that is at least 95 % identical to them,
CC or a sequence having one or more conservative substitutions in them. The
CC polypeptide is useful in manufacturing a medicament for treating a
CC syndrome associated with a human disease selected from a pathology
CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
CC and antibodies are useful in treating or preventing NOVX-associated
CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
CC associated cachexia, and other wasting disorders associated with chronic
CC diseases. The nucleic acids and polypeptides may also be used as targets
CC for the identification of small molecules that modulate or inhibit e.g.
CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
CC wound healing and angiogenesis, in gene therapy, in generation of
CC antibodies that bind immunospecifically to NOVX substances for use in
CC therapeutic or diagnostic methods. The nucleic acids are further used as
CC hybridisation probes, in chromosome mapping, tissue typing, preventive
CC medicine, and pharmacogenomics. The polypeptides are also useful as

CC vaccines. This is the amino acid sequence of a novel human G-protein
CC coupled receptor related protein NOV
XX
SQ Sequence 644 AA;
Query Match 100.0%; Score 660; DB 6; Length 644;
Best Local Similarity 100.0%; Pred. No. 2.3e-65;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLNANNATFYFKIDNVKARQVYV 60
DB 253 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLNANNATFYFKIDNVKARQVYV 312
QY 61 AGKYYFIDFVARETTCSKESNEELTESCTKGLQSLDCNAEYVYVPEKKIYPTVNCQP 120
DB 313 AGKYYFIDFVARETTCSKESNEELTESCTKGLQSLDCNAEYVYVPEKKIYPTVNCQP 372
QY 121 LGM 123
DB 373 LGM 375
RESULT 14
ABU99145
ID ABU99145 standard; protein; 644 AA.
XX AC ABU99145;
XX DT 01-AUG-2003 (first entry)
XX DE Novel human GPCR related protein NOV12c.
XX Human; G-protein coupled receptor related protein; GPCR related protein;
KW NOV; cytostatic; cardiant; antiarteriosclerotic; antidiabetic;
KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
KW diabetes; immune disorder; AIDS; obesity; asthma;
KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
KW infection; multiple sclerosis; cancer-associated cachexia;
KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
XX Homo sapiens.
XX OS
XX WO200299116-A2.
XX PD 12-DEC-2002.
XX 04-JUN-2002; 2002WO-US017428.
XX 04-JUN-2001; 2001US-0295607P.
PR 04-JUN-2001; 2001US-0295661P.
PR 06-JUN-2001; 2001US-0296404P.
PR 06-JUN-2001; 2001US-0296418P.
PR 14-JUN-2001; 2001US-0298285P.
PR 15-JUN-2001; 2001US-0298566P.
PR 21-JUN-2001; 2001US-0299949P.
PR 26-JUN-2001; 2001US-0300883P.
PR 28-JUN-2001; 2001US-0301550P.
PR 13-AUG-2001; 2001US-0311972P.
PR 27-AUG-2001; 2001US-0315071P.
PR 29-AUG-2001; 2001US-0315660P.
PR 14-SEP-2001; 2001US-0322293P.
PR 17-SEP-2001; 2001US-0322706P.
PR 14-DEC-2001; 2001US-0341186P.
PR 17-SEP-2001; 2001US-0322293P.
PR 14-DEC-2001; 2001US-0341186P.
PR 28-FEB-2002; 2002US-0361189P.
PR 12-MAR-2002; 2002US-0363673P.
PR 12-MAR-2002; 2002US-0363676P.
PR 03-JUN-2002; 2002US-00363676.
XX (CURA-) CURAGEN CORP.

XX Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR,
 PI Gangoli EA, Gerlach VL, Gorman L, Guo X, Hjalt T, Kekuda R, Li L,
 PI MacDougall JR, Malyankar UM, Millet I, Padigar M, Patturajan M,
 PI Pena CEA, Rastelli L, Shinkens RA, Stone DJ, Spytek KA, Vernet CM;
 PI Voss EZ, Zernhusen BD;
 XX WPI; 2003-140627/13.
 DR N-PSDB; ACD03649.
 XX
 XX New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 XX Claim 1; Page 144-145; 332pp; English.
 XX
 XX The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 XX Sequence 644 AA;

Query Match 100.0%; Score 660; DB 6; Length 644;
 Best Local Similarity 100.0%; Pred. No. 2.3e-65;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITIKLAENNAATFEKIDNVKARVQV 60
 DB 253 GKDFVQPTKICVGCPRDPTNSPELEETLTHITIKLAENNAATFEKIDNVKARVQV 312
 QY 61 AGKKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 120
 DB 313 AGKKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP 372
 QY 121 LGM 123
 DB 373 LGM 375

RESULT 15
 AAB37447
 ID AAB37447 standard; protein; 122 AA.
 XX
 XX AAB37447;
 XX
 XX 21-FEB-2001 (first entry)
 XX
 XX Human kininogen D3.
 XX
 XX Enzyme; legumain; endopeptidase; cystatin; human; kininogen.
 XX
 XX Homo sapiens.
 XX

FN WC200064945-A1.
 XX
 XX 02-NOV-2000.
 XX
 XX 20-APR-2000; 2000WO-GB001571.
 XX
 XX 22-APR-1999; 99GB-00009133.
 XX
 XX (BABR-) BABRAHAM INST.
 XX
 XX Abrahamson M, Barrett AJ;
 XX WPI; 2000-687316/67.
 XX
 XX Inhibition of mammalian legumain or legumain-related endopeptidase by
 PT cystatin involves interaction with second papain-non-reactive site of
 PT cystatin.
 XX
 XX Disclosure; Fig 4; 45pp; English.
 XX
 XX The present invention relates to inhibition of the enzymatic activity of
 CC legumain or a legumain-related endopeptidase by cystatin. The inhibition
 CC involves an interaction between legumain and a papain-non-reactive site
 CC of cystatin. Legumain (EC 3.4.22.34) is a cysteine endopeptidase, and
 CC performs a protein-processing function. The present sequence is human
 CC kininogen D3, which was used in the present invention. Kininogen is a
 CC type 3 cystatin
 XX
 XX Sequence 122 AA;
 Query Match 95.2%; Score 628; DB 3; Length 122;
 Best Local Similarity 100.0%; Pred. No. 9.5e-63;
 Matches 117; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 7 PPTKICVGCPRDPTNSPELEETLTHITIKLAENNAATFEKIDNVKARVQVAGKKYF 66
 DB 1 PPTKICVGCPRDPTNSPELEETLTHITIKLAENNAATFEKIDNVKARVQVAGKKYF 60
 QY 67 IDFVARETTCKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP LGM 123
 DB 61 IDFVARETTCKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCOP LGM 117

Search completed: September 24, 2004, 14:08:37
 Job time : 52.692 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:04:32 ; Search time 8.364 Seconds
(without alignments)
765.738 Million cell updates/sec

Title: US-10-661-784-1
Perfect score: 660
Sequence: 1 GKDFVQPTKICVGRDIP.....YVPEKKIYPTVNCQPLGM 123

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_42:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	660	100.0	644	1 KNG HUMAN	P01042 homo sapien
2	477	72.3	436	1 KNL1 BOVIN	P01046 bos taurus
3	477	72.3	621	1 KNL1 BOVIN	P01044 bos taurus
4	450	68.2	434	1 KNL2 BOVIN	P01047 bos taurus
5	450	68.2	619	1 KNL2 BOVIN	P01045 bos taurus
6	430	65.2	661	1 KNG MOUSE	O08677 mus musculus
7	426	64.5	639	1 KNG RAT	P08934 rattus norv
8	409	62.0	430	1 KNT2 RAT	P08932 rattus norv
9	401	60.8	430	1 KNT1 RAT	P01048 rattus norv
10	175	26.5	144	1 CYTF MOUSE	O09098 mus musculus
11	165.5	25.1	145	1 CYTF HUMAN	O76096 mus musculus
12	138.5	21.0	149	1 CYTM HUMAN	Q15828 homo sapien
13	136	20.6	148	1 CYTC BOVIN	P01035 bos taurus
14	133.5	20.2	146	1 CYTC MACMU	O19092 macaca mula
15	132.5	20.1	146	1 CYTC SAISC	O19093 saimiri sci
16	130	19.7	127	1 CYTC RAT	P14841 rattus norv
17	129.5	19.6	140	1 CYTC MOUSE	P14460 rattus norv
18	129.5	19.6	378	1 FETB RAT	Q96799 rattus norv
19	129.5	19.6	382	1 FETB HUMAN	Q96799 rattus norv
20	128	19.4	111	1 CYT BITAR	Q96799 rattus norv
21	127.5	19.3	146	1 CYTC HUMAN	P01034 homo sapien
22	125	18.9	141	1 CYTT HUMAN	P09228 homo sapien
23	124.5	18.9	148	1 CYTC RABIT	O07862 oryctolagus
24	122.5	18.6	116	1 CYT COTJA	P10681 coturnix co
25	118.5	18.0	139	1 CYT CHICK	P01038 gallus gall
26	115.5	17.5	388	1 FETB MOUSE	Q96799 rattus norv
27	113.5	17.2	122	1 CYTA SARPE	P11727 sarcophaga
28	113	17.1	141	1 CYTS RAT	P19313 rattus norv
29	109.5	16.6	141	1 CYTN HUMAN	P01037 homo sapien
30	109	16.5	141	1 CYTS HUMAN	P01036 homo sapien
31	108.5	16.4	130	1 CYT ONCKE	Q08967 oncorhynch
32	108	16.4	129	1 CYT CYPCA	P35481 cypripus ca
33	106.5	16.1	130	1 CYT ONCMY	Q01195 oncorhynch

ALIGNMENTS

RESULT 1
KNG_HUMAN
ID KNG_HUMAN STANDARD; PRT; 644 AA.
AC P01042; P01043;
DT 21-JUL-1986 (Rel. 01, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Kininogen precursor (Alpha-2-thiol proteinase inhibitor) [Contains:
DE Bradykinin].
GN KNG.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eureleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
CN NCBI_TaxID=9606;
RX MEDLINE=85234582; PubMed=2989293;
RA Takagaki Y., Kitamura N., Nakanishi S.;
RT "Cloning and sequence analysis of cDNAs for human high molecular
RT weight and low molecular weight prekininogens. Primary structures of
RT two human prekininogens.";
RL J. Biol. Chem. 260:8601-8609(1985).
RN [2]
RP GENE STRUCTURE.
RX MEDLINE=85234583; PubMed=2989294;
RA Kitamura N., Kitagawa H., Fukushima D., Takagaki Y., Miyata T.,
RA Nakanishi S.;
RT "Structural organization of the human kininogen gene and a model for
RT its evolution.";
RL J. Biol. Chem. 260:8610-8617(1985).
RN [3]
RP SEQUENCE OF 1-401 FROM N.A.
RX MEDLINE=85122621; PubMed=6441591;
RA Ohkubo I., Kurachi K., Takasawa T., Shiohara H., Sasaki M.;
RT "Isolation of a human cDNA for alpha 2-thiol proteinase inhibitor and
RT its identity with low molecular weight kininogen.";
RL Biochemistry 23:5691-5697(1984).
RN [4]
RP SEQUENCE OF 379-644.
RX MEDLINE=86030270; PubMed=4054110;
RA Lottspeich F., Kellermann J., Henschen A., Foertsch B.,
RA Mueller-Esterl W.;
RT "The amino acid sequence of the light chain of human high-molecular-
RT mass kininogen.";
RL Eur. J. Biochem. 152:307-314(1985).
RN [5]
RP SEQUENCE OF 381-389.
RX MEDLINE=90255622; PubMed=4952632;
RA Pierce J.V.;
RT "Structural features of plasma kinins and kininogens.";
RL Fed. Proc. 27:52-57(1968).
RN [6]
RP DISULFIDE BONDS.
RA Sueyoshi T., Miyata T., Kato H., Iwanaga S.;
RT "Disulfide bonds in bovine HMW kininogens.";

P22085 onchocerca
Q9d269 mus musculus
Q9h114 homo sapien
P32766 mus musculus
Q9h112 homo sapien
P29699 mus musculus
P81714 naja atra (m
P31726 zea mays (m
P28325 homo sapien
P24090 rattus norv
O60676 homo sapien
O88969 rattus norv

34 105.5 16.0 162 1 CYTX ONCVO
35 101 15.3 139 1 CS11_MOUSE
36 95 14.4 165 1 CS11_HUMAN
37 94.5 14.3 142 1 CST8_MOUSE
38 93.5 14.2 137 1 CS11_HUMAN
39 93.5 14.2 345 1 A2HS_MOUSE
40 93 14.1 99 1 CYT NAJAT
41 91.5 13.9 135 1 CYT1_MAIZE
42 91 13.8 142 1 CYTD_HUMAN
43 90.5 13.7 352 1 A2HS_RAT
44 89.5 13.6 142 1 CST8_HUMAN
45 86.5 13.1 142 1 CST8_RAT

Seikagaku 56:808-808(1984).
 [7]
 RN CARBOHYDRATE-LINKAGE SITE ASN-294.
 RX MEDLINE=22660472; PubMed=12754519;
 RA Zhang H., Li X.-J., Martin D.B., Aebbersold R.;
 RT "Identification and quantification of N-linked glycoproteins using
 hydrazide chemistry, stable isotope labeling and mass spectrometry.";
 RL Nat. Biotechnol. 21:660-666(2003).
 CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 CC HMW-kininogen plays an important role in blood coagulation by
 CC helping to position optimally prekallikrein and factor XI next to
 CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
 CC induced aggregation of thrombocytes; (4) the active peptide
 CC bradykinin that is released from HMW-kininogen shows a variety of
 CC physiological effects: (4A) influence in smooth muscle
 CC contraction, (4B) induction of hypotension, (4C) natriuresis and
 CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
 CC mediator of inflammation and causes (4E1) increase in vascular
 CC permeability, (4E2) stimulation of nociceptors (4E3) release of
 CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
 CC a cardioprotective effect (directly via bradykinin action,
 CC indirectly via endothelium-derived relaxing factor action); (5)
 CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
 CC kininogen is in contrast to HMW-kininogen not involved in blood
 CC clotting.
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- ALTERNATIVE PRODUCTS.
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=HMW;
 CC IsoId=P01042-1; Sequences=Displayed;
 CC Name=LMW;
 CC IsoId=P01042-2; Sequence=VSP_001261, VSP_001262;
 CC -!- TISSUE SPECIFICITY: Plasma.
 CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -!- SIMILARITY: Contains 3 cystatin-like domains.
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 CC or send an email to license@isb-sib.ch).
 CC
 DR EMBL; K02566; AAA35497.1; -;
 DR EMBL; M11437; AAB59550.1; JOINED.
 DR EMBL; M11438; AAB59550.1; JOINED.
 DR EMBL; M11521; AAB59550.1; JOINED.
 DR EMBL; M11522; AAB59550.1; JOINED.
 DR EMBL; M11523; AAB59550.1; JOINED.
 DR EMBL; M11524; AAB59550.1; JOINED.
 DR EMBL; M11525; AAB59550.1; JOINED.
 DR EMBL; M11526; AAB59550.1; JOINED.
 DR EMBL; M11527; AAB59550.1; JOINED.
 DR EMBL; M11528; AAB59550.1; JOINED.
 DR EMBL; M11529; AAB59550.1; JOINED.
 DR EMBL; M11437; AAB59551.1; -;
 DR EMBL; M11438; AAB59551.1; JOINED.
 DR EMBL; M11521; AAB59551.1; JOINED.
 DR EMBL; M11522; AAB59551.1; JOINED.
 DR EMBL; M11523; AAB59551.1; JOINED.
 DR EMBL; M11524; AAB59551.1; JOINED.
 DR EMBL; M11525; AAB59551.1; JOINED.
 DR EMBL; M11526; AAB59551.1; JOINED.
 DR EMBL; M11527; AAB59551.1; JOINED.
 DR EMBL; M11528; AAB59551.1; JOINED.
 DR PIR; A01279; KGHU1.
 DR PIR; A01280; KGHU1.
 DR SWISS-2DPAGE; P01042; HUMAN.
 DR Genew; HGNC:6383; KNG.
 DR MIM; 228960; -;
 DR GO; GO:0007596; P:blood coagulation; NAS.
 DR GO; GO:0030146; P:diuresis; NAS.
 DR GO; GO:0006954; P:inflammatory response; NAS.

DR GO; GO:0030147; P:natriuresis; NAS.
 DR GO; GO:0006939; P:smooth muscle contraction; NAS.
 DR InterPro; IPR000010; Cystatin.
 DR InterPro; IPR002395; Kininogen.
 DR Pfam; PF00031; cystatin; 3.
 DR PRINTS; PR000314; KININOGEN.
 DR SMART; SM00043; CY; 3.
 DR PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Thiol protease inhibitor; Vasodilator;
 KW Bradykinin; Blood coagulation; Inflammatory response; Signal;
 KW Alternative splicing; Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 644 KININOGEN.
 FT CHAIN 19 380 KININOGEN HEAVY CHAIN.
 FT PEPTIDE 381 389 BRADYKININ.
 FT CHAIN 390 644 KININOGEN LIGHT CHAIN.
 FT DOMAIN 19 136 CYSTATIN-LIKE 1.
 FT DOMAIN 137 258 CYSTATIN-LIKE 2.
 FT DOMAIN 259 380 CYSTATIN-LIKE 3.
 FT DOMAIN 420 510 HIS-RICH
 FT (ASSOCIATED WITH CLOTTING ACTIVITY).
 FT REPEAT 420 449
 FT REPEAT 450 479
 FT REPEAT 480 510
 FT MOD_RES 19 19
 FT DISULFID 28 614
 FT DISULFID 83 94
 FT DISULFID 107 126
 FT DISULFID 142 145
 FT DISULFID 206 218
 FT DISULFID 229 248
 FT DISULFID 264 267
 FT DISULFID 328 340
 FT DISULFID 351 370
 FT CARBOHYD 48 48 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 169 169 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 205 205 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 294 294 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 401 401 O-LINKED.
 FT CARBOHYD 533 533 O-LINKED.
 FT CARBOHYD 542 542 O-LINKED.
 FT CARBOHYD 546 546 O-LINKED.
 FT CARBOHYD 557 557 O-LINKED.
 FT CARBOHYD 571 571 O-LINKED.
 FT CARBOHYD 577 577 O-LINKED.
 FT CARBOHYD 593 593 O-LINKED.
 FT CARBOHYD 628 628 O-LINKED.
 FT VARSPLIC 402 427 VSPHTSMAPAQDEERDSGKQCHTR -> SHLRSCYKGR
 PKGAEPASEREVS (in isoform LMW).
 /FTId-VSP_001261.
 Missing (in isoform LMW).
 /FTId-VSP_001262.
 T -> I (IN REF. 1).
 Query Match 100.0%; Score 660; DB 1; Length 644;
 Best Local Similarity 100.0%; Pred. No. 2e-55; 0; Gaps 0;
 Matches 123; Conservative 0; Mismatches 0; Indels 0;
 Qy 1 GKDFVQPTKICVGCPRDIPTNSPELEETLTHITKLNANNATFYFKIDNVKARVQV 60
 Db 253 GKDFVQPTKICVGCPRDIPTNSPELEETLTHITKLNANNATFYFKIDNVKARVQV 312
 Qy 61 AGKYSIDFVARETTCKSKESNELTSCCKLQSLDCNAEYVVPWEKKIYPTVNCOP 120
 Db 313 AGKYSIDFVARETTCKSKESNELTSCCKLQSLDCNAEYVVPWEKKIYPTVNCOP 372
 Qy 121 LGM 123
 Db 373 LGM 375
 RESULT 2

KNL1_BOVIN
ID KNL1_BOVIN STANDARD; PRT; 436 AA.
AC P01046;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Kininogen, LMW I precursor (Thiol proteinase inhibitor) [Contains:
Bradykinin].
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=83117859; PubMed=6572010;
RA Nawa H., Kitamura N., Hirose T., Asai M., Inayama S., Nakanishi S.;
RT "Primary structures of bovine liver low molecular weight kininogen
precursors and their two mRNAs.";
RL Proc. Natl. Acad. Sci. U.S.A. 80:90-94 (1983).
RN [2]
RP SEQUENCE OF 19-378.
RX MEDLINE=87137530; PubMed=3546295;
RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
RA Miyata T., Iwanaga S.;
RT "Bovine high molecular weight kininogen. The amino acid sequence,
positions of carbohydrate chains and disulfide bridges in the heavy
chain portion.";
RL J. Biol. Chem. 262:2768-2779 (1987).
CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
LMW-kininogen inhibits the aggregation of thrombocytes; (3) the
active peptide kallidin that is released from LMW-kininogen shows
a variety of physiological effects: (3A) influence in smooth
muscle contraction, (3B) induction of hypotension, (3C)
natriuresis and diuresis (kidney).
CC -!- SUBCELLULAR LOCATION: Extracellular.
CC -!- ALTERNATIVE PRODUCTS:
Event=Alternative splicing; Named isoforms=2;
Name=LMW I;
CC Name=HMW I;
CC IsoId=P01046-1; Sequence=Displayed;
CC IsoId=P01044-1; Sequence=External;
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -!- MISCELLANEOUS: LMW-kininogen is in contrast to HMW-kininogen not
involved in blood clotting.
CC -!- SIMILARITY: Contains 3 cystatin-like domains.

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or send an email to license@isb-sib.ch).

DR EMBL; V00426; CAA23709.1; -;
DR PIR; A01283; KGBOL1.
DR InterPro; IP3000010; Cystatin.
DR Pfam; PF00031; cystatin; 3.
DR SMART; SM00043; Cy; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
KW Thiol protease inhibitor; Bradykinin; Signal;
KW Pyrrolidone carboxylic acid.
FT CHAIN 1 18 KININOGEN, LMW I.
FT CHAIN 19 436 HEAVY CHAIN.
FT PEPTIDE 19 378 BRADYKININ.
FT CHAIN 380 388 LIGHT CHAIN.
FT DOMAIN 389 436 CYPSTATIN-LIKE 1.
FT DOMAIN 19 135 CYPSTATIN-LIKE 2.
FT DOMAIN 136 257 CYPSTATIN-LIKE 3.
FT DOMAIN 258 378 CYPSTATIN-LIKE 3.

MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
FT CARBOHYD 87 87 N-LINKED (GLCNAC. . .).
FT CARBOHYD 136 136 O-LINKED (PARTIAL. . .).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
FT DISULFID 27 406 INTERCHAIN.
FT DISULFID 82 93
FT DISULFID 106 125
FT DISULFID 141 144
FT DISULFID 205 217
FT DISULFID 228 247
FT DISULFID 263 286
FT DISULFID 327 339
FT DISULFID 350 369
FT CONFLICT 295 295 A -> T (IN REF. 1; CAA23709).
SQ SEQUENCE 436 AA; 48427 MW; F01F7EB6814BCE6C CRC64;
Query Match 72.3%; Score 477; DB 1; Length 436;
Best Local Similarity 71.9%; Pred. No. 3.6e-38;
Matches 87; Conservative 14; Mismatches 20; Indels 0; Gaps 0;
QY 2 KDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNATFYFKIDNVKARQVVA 61
DB 253 KDFVQPTKLCAGCPKIPVDSFDLEELSHSIAKLAHEDGAFYFKIDTVKATQVVA 312
QY 62 GKRYFIDFVARETTCSENEELTSCETPKLGQSLDCNAEYVYVWPEKKIYPTVNCQPL 121
DB 313 GLKYSIVFIARETTCSENEELTKSGNEELTKSCINIHQILHCDANVYVWPEKKYPTVNCQPL 372
QY 122 G 122
DB 373 G 373
RESULT 3
KNL1_BOVIN STANDARD; PRT; 621 AA.
ID KNL1_BOVIN
AC P01044;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Kininogen, HMW I precursor (Thiol proteinase inhibitor) [Contains:
Bradykinin].
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=84014106; PubMed=6571699;
RA Kitamura N., Takagaki Y., Furuto S., Tanaka T., Nawa H., Nakanishi S.;
RT "A single gene for bovine high molecular weight and low molecular
weight kininogens.";
RL Nature 305:545-549 (1983).
RN [2]
RP SEQUENCE OF 19-378.
RX MEDLINE=87137530; PubMed=3546295;
RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
RA Miyata T., Iwanaga S.;
RT "Bovine high molecular weight kininogen. The amino acid sequence,
positions of carbohydrate chains and disulfide bridges in the heavy
chain portion.";
RL J. Biol. Chem. 262:2768-2779 (1987).
RN [3]
RP SEQUENCE OF 378-393.
RX MEDLINE=70180420; PubMed=4986212;
RA Kato H., Nagasawa S., Suzuki T.;
RT "Studies on the structure of bovine kininogen: cleavages of disulfide
bonds and of methionyl bonds in kininogen-II.";
RL J. Biochem. 67:313-323 (1970).
RN [4]

RP SEQUENCE OF 458-498.
RX MEDLINE=75170265; PubMed=1169237;
RA Han Y N., Komiya M., Iwanaga S., Suzuki T.;
RT "Studies on the primary structure of bovine high-molecular-weight
RT kininogen. Amino acid sequence of a fragment ('histidine-rich
RT peptide') released by plasma kallikrein."
RL J. Biochem. 77:55-68(1975).
CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC HMW-kininogen plays an important role in blood coagulation by
CC helping to position optimally prekallikrein and factor XI next to
CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
CC induced aggregation of thrombocytes; (4) the active peptide
CC bradykinin that is released from HMW-kininogen shows a variety of
CC physiological effects: (4A) influence in smooth muscle
CC contraction, (4B) induction of hypotension, (4C) natriuresis and
CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
CC mediator of inflammation and causes (4E1) increase in vascular
CC permeability, (4E2) stimulation of nociceptors (4E3) release of
CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action).
CC -!- SUBCELLULAR LOCATION: Extracellular.
CC -!- ALTERNATIVE PRODUCTS:
CC Event-Alternative splicing; Named isoforms=2;
CC Name=HMW I;
CC IsoId=P01044-1; Sequences=Displayed;
CC Name=LMW I;
CC IsoId=P01046-1; Sequences=External;
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -!- SIMILARITY: Contains 3 cystatin-like domains.
CC
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CC
CC -----
CC EMBL: V01491; CAA24735.1; -
CC PIR: A01281; KGBOH1.
CC InterPro: IPR000010; Cystatin.
CC InterPro: IPR002395; Kininogen.
CC Pfam: PF00031; Cystatin_3.
CC PRINTS: PRO0334; KININOGEN.
CC SMART: SM00043; CY; 3.
CC PROSITE: PS00287; CYSTATIN; 2.
CC Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
CC Thiol protease inhibitor; Bradykinin; Blood coagulation;
CC Inflammatory response; Signal; Pyrrolidone carboxylic acid.
CC SIGNAL 1 18 PROBABLE
CC CHAIN 19 621 KININOGEN, HMW I.
CC CHAIN 19 378 HEAVY CHAIN.
CC PEPTIDE 380 388 BRADYKININ.
CC CHAIN 389 621 LIGHT CHAIN.
CC DOMAIN 19 135 CYSTATIN-LIKE 1.
CC DOMAIN 136 257 CYSTATIN-LIKE 2.
CC DOMAIN 258 378 CYSTATIN-LIKE 3.
CC MOD_RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
CC CARBOHYD 87 87 N-LINKED (GLCNAC. . .).
CC CARBOHYD 136 136 O-LINKED (PARTIAL).
CC CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
CC CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
CC CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
CC DISULFID 27 591 INTERCHAIN.
CC DISULFID 82 93
CC DISULFID 106 125
CC DISULFID 141 144
CC DISULFID 205 217
CC DISULFID 228 247
CC DISULFID 263 266
CC DISULFID 327 339

FT DISULFID 350 369
SQ SEQUENCE 621 AA; 68890 MW; D16850BEFE3C55CD CRC64;
Query Match 72.3%; Score 477; DB 1; Length 621;
Best Local Similarity 71.9%; Pred. No. 5.4e-38;
Matches 87; Conservative 14; Mismatches 20; Indels 0; Gaps 0;
QY 2 KDFVQPTKICVGCPRDIPNSPELEETLTHITKLAENNAFFKIDNVKARVQVVA 61
DB 253 KDFVQPTKICVGCPRDIPNSPELEETLTHITKLAENNAFFKIDNVKARVQVVA 312
QY 62 GKQYFDVARETCKSKESNEELTSCETKLGQSLDCNAEYVVPWEKIVPTVNCQPL 121
DB 313 GLKYSIVFIARETCKSGSNEELTSCSEININGQILHCDANVYVVPWEKIVPTVNCQPL 372
QY 122 G 122
DB 373 G 373
RESULT 4
KML2 BOVIN STANDARD; PRT; 434 AA.
ID KML2 BOVIN
AC P01047;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Kininogen, LMW II precursor (Thiol proteinase inhibitor) [Contains:
DE Bradykinin].
DE Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OC NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=83117859; PubMed=6572010;
RA Nawa H., Kitamura N., Hirose T., Asai M., Inayama S., Nakanishi S.;
RT "Primary structures of bovine liver low molecular weight kininogen
RT precursors and their two mRNAs".
RL Proc. Natl. Acad. Sci. U.S.A. 80:90-94 (1983).
RN [2]
RP SEQUENCE OF 19-376..
RX MEDLINE=87137530; PubMed=3546285;
RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
RA Miyata T., Iwanaga S.;
RT "Bovine high molecular weight kininogen. The amino acid sequence,
RT positions of carbohydrate chains and disulfide bridges in the heavy
RT chain portion."
RL J. Biol. Chem. 262:2768-2779 (1987).
CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC LMW-kininogen inhibits the aggregation of thrombocytes; (3) the
CC active peptide kallidin that is released from LMW-kininogen shows
CC a variety of physiological effects: (3A) influence in smooth
CC muscle contraction, (3B) induction of hypotension, (3C)
CC natriuresis and diuresis (kidney).
CC -!- SUBCELLULAR LOCATION: Extracellular.
CC -!- ALTERNATIVE PRODUCTS:
CC Event-Alternative splicing; Named isoforms=2;
CC Name=LMW II;
CC IsoId=P01047-1; Sequence=Displayed;
CC Name=HMW II;
CC IsoId=P01045-1; Sequence=External;
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -!- MISCELLANEOUS: LMW-kininogen is in contrast to HMW-kininogen not
CC involved in blood clotting.
CC -!- SIMILARITY: Contains 3 cystatin-like domains.
CC
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DR EMBL; V00427; CAA23710.1; -.
DR PIR; A01284; KGBOL2.
DR HSP; P01038; IA90.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 3.
DR SMART; SM00043; Cy; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Vasodilator; Alternative splicing;
KW Thiol protease inhibitor; Bradykinin; Signal;
KW Pyrrolidone carboxylic acid.
FT SIGNAL 1
FT CHAIN 19 434 KININOGEN, LMW II.
FT CHAIN 19 376 HEAVY CHAIN.
FT PEPTIDE 378 386 BRADYKININ.
FT CHAIN 387 434 LIGHT CHAIN.
FT DOVAIN 19 135 CYSTATIN-LIKE 1.
FT DOVAIN 136 256 CYSTATIN-LIKE 2.
FT DOVAIN 257 376 CYSTATIN-LIKE 3.
FT MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
FT CARBOHYD 87 87 O-LINKED (GLCNAC. . .).
FT CARBOHYD 136 136 O-LINKED (PARTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
FT CARBOHYD 280 280 N-LINKED (GLCNAC. . .).
FT DISULFID 27 404 INTERCHAIN.
FT DISULFID 82 93
FT DISULFID 106 125
FT DISULFID 141 144
FT DISULFID 205 217
FT DISULFID 228 247
FT DISULFID 261 264
FT DISULFID 325 337
FT DISULFID 348 367
SQ SEQUENCE 434 AA; 48148 MW; 73A7079DB3E03430 CRC64;

Query Match 68.2%; Score 450; DB 1; Length 434;
Best Local Similarity 68.9%; Pred. No. 1.4e-35;
Matches 84; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

QY 1 GKDFVQPTKICVGPDRIPNTSPLEBTLTHITKLNAENNAFYKIDNVKKARQVQV 60
Db 252 GEDEL--PPMVCVCPKPIVDSPDLSEALNHSIAKLNAEHDGFYKIDTVKKATQVQV 309
QY 61 AGKXYFIDFVARETTCSEKSENEELTESCETKKLQSLDCNAEVVYVPWEKKIYPTVNCQP 120
Db 310 GGLKYSIVFIARETTCSEKSENEELTKSCEINIHQILHCDANVYVPWEKKVYPTVNCQP 369
QY 121 LG 122
Db 370 LG 371

RESULT 5
KNH2_BOVIN
ID_KNH2_BOVIN STANDARD; PRT; 619 AA.
AC P01045;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Kininogen, HMW II precursor (Thiol proteinase inhibitor) [Contains:
DE Bradykinin].
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.

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RX MEDLINE=84014106; PubMed=6571699;
RA Kitamura N., Takaaki Y., Furuto S., Tanaka T., Nawa H., Nakanishi S.;
RT "A single gene for bovine high molecular weight and low molecular
RT weight kininogens";
RL Nature 305:545-549(1983).
RN [2]
RP SEQUENCE OF 19-376.
RX MEDLINE=87137530; PubMed=3546295;
RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
RA Miyata T., Iwanaga S.;
RT "Bovine high molecular weight kininogen. The amino acid sequence,
RT positions of carbohydrate chains and disulfide bridges in the heavy
RT chain portion.";
RL J. Biol. Chem. 262:2768-2779(1987).
RN [3]
RP SEQUENCE OF 376-391.
RX MEDLINE=70180420; PubMed=4986212;
RA Kato H., Nagasawa S., Suzuki T.;
RT "Studies on the structure of bovine kininogen: cleavages of disulfide
RT bonds and of methionyl bonds in kininogen-II.";
RL J. Biochem. 67:313-323(1970).
RN [4]
RP SEQUENCE OF 387-455.
RX MEDLINE=76260155; PubMed=956151;
RA Han Y.N., Kato H., Iwanaga S., Suzuki T.;
RT "Primary structure of bovine plasma high-molecular-weight kininogen.
RT The amino acid sequence of a glycopeptide portion (fragment 1)
RT following the C-terminus of the bradykinin moiety.";
RL J. Biochem. 77:55-68(1975).
RN [5]
RP SEQUENCE OF 456-496.
RX MEDLINE=75170265; PubMed=1169237;
RA Han Y.N., Komiya M., Iwanaga S., Suzuki T.;
RT "Studies on the primary structure of bovine high-molecular-weight
RT kininogen. Amino acid sequence of a fragment ('histidine-rich
RT peptide') released by plasma kallikrein.";
RL J. Biochem. 77:55-68(1975).
CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC HMW-kininogen plays an important role in blood coagulation by
CC helping to position optimally prekallikrein and factor XI next to
CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
CC induced aggregation of thrombocytes; (4) the active peptide
CC bradykinin that is released from HMW-kininogen shows a variety of
CC physiological effects: (4A) influence in smooth muscle
CC contraction, (4B) induction of hypotension, (4C) natriuresis and
CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
CC mediator of inflammation and causes (4E1) increase in vascular
CC permeability, (4E2) stimulation of nociceptors (4E3) release of
CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action).
CC -!- SUBCELLULAR LOCATION: Extracellular.
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=HMW II;
CC IsoId=P01045-1; Sequence=Displayed;
CC Name=LMW II;
CC IsoId=P01047-1; Sequence=External;
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -!- SIMILARITY: Contains 3 cystatin-like domains.
CC -----
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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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DR InterPro: IPR000010; Cystatin.
DR InterPro: IPR002395; Kininogen.
DR Pfam: PF00031; Cystatin_3.
DR PRINTS: PRO0334; KININOGEN.
DR SMART: SM00043; CY; 3.
DR PROSITE: PS00287; CYSTATIN; 2.
DR GlycoProtein: Plasma; Repeat; Vasodilator; Alternative splicing;
KW Thiol protease inhibitor; Bradykinin; Blood coagulation; Signal;
KW Inflammatory response; Pyrrolidone carboxylic acid.
PT SIGNAL 1 18
PT CHAIN 19 619 KININOGEN, HWY II.
PT PEPTIDE 378 386 HEAVY CHAIN.
PT CHAIN 387 619 BRADYKININ.
PT CHAIN 387 619 LIGHT CHAIN.
PT DOMAIN 19 135 CYSTATIN-LIKE 1.
PT DOMAIN 136 256 CYSTATIN-LIKE 2.
PT DOMAIN 257 376 CYSTATIN-LIKE 3.
PT MOD_RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
PT CARBOHYD 87 97 N-LINKED (GLCNAC. . .).
PT CARBOHYD 136 136 O-LINKED (PARTIAL. . .).
PT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
PT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
PT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
PT CARBOHYD 280 280 N-LINKED (GLCNAC. . .).
PT CARBOHYD 400 400 O-LINKED.
PT DISULFID 27 589 INTERCHAIN.
PT DISULFID 82 93
PT DISULFID 105 125
PT DISULFID 141 144
PT DISULFID 205 217
PT DISULFID 228 247
PT DISULFID 261 284
PT DISULFID 325 337
PT DISULFID 348 367
PT VARIANT 398 398
PT VARIANT 401 401
PT VARIANT 454 454
PT VARIANT 454 454
SQ SEQUENCE 619 AA; 68710 MW; F04320A8EB0E50DA CRC64;

Query Match 68.2%; Score 450; DB 1; Length 619;
Best Local Similarity 68.9%; Pred. No. 2e-35;
Matches 84; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

QY 1 GKDVPQPKICVGPDRDPTNSPLEETLTITIKLAENNATFYKIDNVKKARVQV 60
Db 252 GDFL--PMWCVGCPKIPVDSPDEALNHSIAKLAHDTGYFKIDTVKKATVQV 309

QY 61 AGKKYFIDFVARETTCKSESNEELTESCTKKLGSLDPCNAFVYVPWEKKIYPTVNCQ 120
Db 310 GGLKYSIVFIARETTCKSGSNEELTKSCBINHTGQILHCDANVYVPWEKKYPTVNCQ 369

QY 121 LG 122
Db 370 LG 371

RESULT 6
ID KNG MOUSE STANDARD; PRT; 661 AA.
AC O08677; O08676; Q91XK5;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Kininogen precursor [Contains: Bradykinin].
GN KNG.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORMS HWY AND LMW).
RC STRAIN=C57BL/6 X CBA; TISSUE=Liver;
RC MEDLINE=97342556; PubMed=9199253;

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RA Takano M., Kondo J., Yayama K., Otani M., Sano K., Okamoto H.;
RT "Molecular cloning of cDNAs for mouse low-molecular-weight and high-
RT molecular-weight prekininogens.";
RL Biochim. Biophys. Acta 1352:222-230(1997).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORM LMW).
RC STRAIN=C57BL/6J; TISSUE=Placenta;
RX MEDLINE=22354683; PubMed=1246851;
RA Okazaki Y., Furuno M., Kasukawa T., Adachi H., Yamanaka I., Kiyosawa H.,
RA Nikaido I., Osato N., Saito R., Suzuki H., Schonbach C., Gojobori T.,
RA Yagi K., Tomaru Y., Hasegawa Y., Nogami A., Quackenbush J.,
RA Baldarelli R., Hill D.P., Bult C., Hume D.A., Quackenbush J.,
RA Schriml L.M., Karspin A., Matsuda H., Batalov S., Beisel K.W.,
RA Blake J.A., Bratt D., Brusic V., Chothia C., Corbani L.E., Cousins S.,
RA Dalla E., Dragani T.A., Fletcher C.F., Forrest A., Frazer K.S.,
RA Gaasterland T., Gariboldi M., Gissi C., Godzik A., Gough J.,
RA Grimmond S., Gustincich S., Hirokawa N., Jackson I.J., Jarvis E.D.,
RA Kanai A., Kawaji H., Kawasawa Y., Kedzierski R.M., King B.L.,
RA Konagaya A., Kurochkin I.V., Lee Y., Lenhard B., Lyons P.A.,
RA Maglott D.R., Maltais L., Marchionni L., McKenzie L., Miki H.,
RA Nagashima T., Numata K., Okido T., Pavan W.J., Perlea G., Pesole G.,
RA Petrovsky N., Pillai R., Pontius J.U., Qi D., Ramachandran S.,
RA Ravasi T., Reed J.C., Reed D.J., Reid J., Ring B.Z., Ringwald M.,
RA Sandelin A., Schneider C., Semple C.A., Setou M., Shimada K.,
RA Sultana R., Takenaka Y., Taylor M.S., Teasdale R.D., Tomita M.,
RA Verardo R., Wagner L., Wahlestedt C., Wang Y., Watanabe Y., Wells C.,
RA Wilming L.G., Wynshaw-Boris A., Yanagisawa M., Yang I., Yang L.,
RA Yuan Z., Zavolan M., Zhu Y., Zimmer A., Carninci P., Hayatsu N.,
RA Hirozane-Kishikawa T., Konno H., Nakamura M., Sakazume N., Sato K.,
RA Shiraki T., Waki K., Kawai J., Aizawa K., Arakawa T., Fukuda S.,
RA Hara A., Hashizume W., Imotani K., Ishii Y., Itoh M., Kagawa I.,
RA Miyazaki A., Sakai K., Sasaki D., Shibata K., Shinagawa A.,
RA Yasunishi A., Yoshino M., Waterston R., Lander E.S., Rogers J.,
RA Birney E., Hayashizaki Y.;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs.";
RL Nature 420:563-573(2002).
RN [3]
RP SEQUENCE FROM N.A. (ISOFORM LMW).
RC TISSUE=Liver;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G., Schuler G.D.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Scapleton M., Soares M.B., Donald M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Udell T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
RA Whitting M., Madan A., Touchman J.W., Green E.D., Dickson M.C.,
RA Blakesley R.W., Touchman J.W., Schmutz J., Myers R.M.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Skalska U., Smalish D.E.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalish D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC HWY-kininogen plays an important role in blood coagulation by
CC helping to position optimally prekallikrein and factor XI next to
CC factor XII; (3) HWY-kininogen inhibits the thrombin and plasmin-
CC induced aggregation of thrombocytes; (4) the active peptide
CC bradykinin that is released from HWY-kininogen shows a variety of
CC physiological effects: (4A) influence in smooth muscle
CC contraction, (4B) induction of hypotension, (4C) patriuresis and
CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
CC mediator of inflammation and causes (4E1) increase in vascular
CC permeability, (4E2) stimulation of nociceptors (4E3) release of
CC other mediators of inflammation (e.g. prostaglandins), (4F) it has

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CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action); (5)
CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
CC kininogen is in contrast to HMW-kininogen not involved in blood
CC clotting (By similarity).
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=HMW;
CC IsoId=008677-1; Sequence=Displayed;
CC Name=LMW;
CC IsoId=008677-2; Sequence=VSP_001263, VSP_001264;
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- PTM: Bradykinin is released from kininogen by plasma kallikrein.
CC -!- SIMILARITY: Contains 3 cysteine-like domains.
CC -----
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CC -----
DR EMBL; D84435; BAA19743.1; -;
DR EMBL; D84415; BAA19742.1; -;
DR EMBL; AK005547; BAB24115.1; -;
DR EMBL; BC018158; AAH18158.1; -;
DR MGB; MGI:1097705; Kug.
DR InterPro; IPR000010; Cystatin.
DR InterPro; IPR002395; Kininogen.
DR Pfam; PF00031; cystatin; 3.
DR PRINTS; PR00334; KININOGEN.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 1.
KW Glycoprotein, Plasma; Repeat; Thiol protease inhibitor; Vasodilator;
KW Bradykinin; Blood coagulation; Inflammatory response; Signal;
KW Alternative splicing.
FT SIGNAL 1 18
FT CHAIN 19 661 KININOGEN.
FT CHAIN 19 379 KININOGEN HEAVY CHAIN.
FT CHAIN 380 388 BRADYKININ.
FT CHAIN 389 661 KININOGEN LIGHT CHAIN.
FT DOMAIN 19 135 CYSTATIN-LIKE 1.
FT DOMAIN 136 257 CYSTATIN-LIKE 2.
FT DOMAIN 258 379 CYSTATIN-LIKE 3.
FT DOMAIN 439 524 HIS-RICH.
FT DISULFD 28 631 INTERCHAIN (BY SIMILARITY).
FT DISULFD 83 94 BY SIMILARITY.
FT DISULFD 107 125 BY SIMILARITY.
FT DISULFD 141 144 BY SIMILARITY.
FT DISULFD 205 217 BY SIMILARITY.
FT DISULFD 228 247 BY SIMILARITY.
FT DISULFD 263 266 BY SIMILARITY.
FT DISULFD 327 339 BY SIMILARITY.
FT DISULFD 350 369 BY SIMILARITY.
FT CARBOHYD 82 82 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 242 242 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT VARSPIC 401 432 VSPPIAREQERDAETEQGTHGWLHEKO -> RLIRA
FT CEYGRLSKAGAEPAPEQAESSQVKQ (in isoform
FT /FTId=VSP_001263.
FT Missing (in isoform LMW).
FT /FTId=VSP_001264.
SQ SEQUENCE 661 AA; 73102 MW; 774460258D58796E CRC64;
Query Match 65.2%; Score 430; DB 1; Length 661;
Best Local Similarity 65.9%; Pred. No. 1.8e-33;
Matches 81; Conservative 11; Mismatches 33; Indels 0; Gaps 0;
QY 1 GKDFVQPTKICVCGPRDIPNSPELEBTLTHITKLAENNAATFYFKIDNVKARQVQV 60

Db 252 GDVLVAPLKPFGCPDRDIPVDSPELKVGLHSIAQLNAENHPFYKIDTVKATSQV 311
QY 61 AGKXYFIDFVARETCKESNEELTESCETKLQSLDCNARVYVVPVKKYIPTVNCOP 120
Db 312 AGTKYVIEFIARETKCKESNTELAEDCEIKHLQSLDCNARVYVVPVKKYIPTVNCOP 371
QY 121 LGM 123
Db 372 LDM 374
RESULT 7
KNG_RAT
ID_KNG_RAT STANDARD; PRT; 639 AA.
AC P08934; P08933;
DT 01-NOV-1988 (Rel. 09, Created)
DT 01-NOV-1988 (Rel. 09, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Kininogen precursor [Contains: Bradykinin].
GN KNG.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP MEDLINE=87137443; PubMed=3029068;
RA Kitagawa H., Kitamura N., Hayashida H., Miyata T., Nakanishi S.;
RT "Differing expression patterns and evolution of the rat kininogen
RT gene family.";
RL J. Biol. Chem. 262:2190-2198(1987).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORMS HMW AND LMW).
RA Puroto-Rato S., Matsumoto A., Kitamura N., Nakanishi S.;
RT "Primary structures of the mRNAs encoding the rat precursors for
RT bradykinin and T-kinin. Structural relationship of kininogens with
RT major acute phase protein and alpha 1-cysteine proteinase
RT inhibitor.";
RL J. Biol. Chem. 260:12054-12059(1985).
RN [3]
RP SEQUENCE OF 1-65 FROM N.A.
RC STRAIN=Buffalo;
RX MEDLINE=87250580; PubMed=2439509;
RA Fung W.-P., Schreiber G.;
RT "Structure and expression of the genes for major acute phase alpha 1-
RT protein (thiostratin) and kininogen in the rat.";
RL J. Biol. Chem. 262:9298-9308(1987).
RN [4]
RP SEQUENCE OF 1-41 FROM N.A.
RC STRAIN=Wistar; TISSUE=Liver;
RX MEDLINE=87137465; PubMed=3818598;
RA Kageyama R., Kitamura N., Ohkubo H., Nakanishi S.;
RT "Differing utilization of homologous transcription initiation sites
RT of rat K and T kininogen genes under inflammation condition.";
RL J. Biol. Chem. 262:2345-2351(1987).
CC -!- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
CC HMW-kininogen plays an important role in blood coagulation by
CC helping to position optimally prekallikrein and factor XI next to
CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
CC induced aggregation of thrombocytes; (4) the active peptide
CC bradykinin that is released from HMW-kininogen shows a variety of
CC physiological effects: (4A) influence in smooth muscle
CC contraction, (4B) induction of hypotension, (4C) natriuresis and
CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
CC mediator of inflammation and causes (4E1) increase in vascular
CC permeability, (4E2) stimulation of nociceptors (4E3) release of
CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
CC a cardioprotective effect (directly via bradykinin action,
CC indirectly via endothelium-derived relaxing factor action); (5)
CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
CC kininogen is in contrast to HMW-kininogen not involved in blood

DR PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;
 KW Thiol protease inhibitor; Bradykinin; Acute phase; Signal.
 FT SIGNAL 1 18
 FT CHAIN 19 430 KININOGEN, T-II.
 FT CHAIN 19 375 HEAVY CHAIN.
 FT PEPTIDE 376 386 T-KININ.
 FT CHAIN 387 430 LIGHT CHAIN.
 FT DOMAIN 19 135 CYSTATIN-LIKE 1.
 FT DOMAIN 136 257 CYSTATIN-LIKE 2.
 FT DOMAIN 258 375 CYSTATIN-LIKE 3.
 FT DOMAIN 28 404 INTERCHAIN (BY SIMILARITY).
 FT DISULFID 83 94 BY SIMILARITY.
 FT DISULFID 107 125 BY SIMILARITY.
 FT DISULFID 208 217 BY SIMILARITY.
 FT DISULFID 228 247 BY SIMILARITY.
 FT DISULFID 263 266 BY SIMILARITY.
 FT DISULFID 327 339 BY SIMILARITY.
 FT DISULFID 350 369 BY SIMILARITY.
 FT CARBOHYD 82 82 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 126 126 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 326 326 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 430 AA; 47524 MW; 43EDF02D1BF55076 CRC64;
 Query Match 62.08; Score 409; DB 1; Length 430;
 Best Local Similarity 61.8%; Pred. No. 1.1e-31;
 Matches 76; Conservative 15; Mismatches 32; Indels 0; Gaps 0;
 QY 1 GKDFVQPTKICVGPDIPTNSPELEETLTHITIKLAENNAATFFPKIDNVKARVQVY 60
 DB 252 GDDLFSLLPKFCGCPKPNIPVDSPELKEALGHSLAQLNAQHNLFFPKIDTVKATSQVY 311
 QY 61 AGKYFIDFVAREFTCSKESEBELTESCEKLGQSLDCAEYVVPWEKKIYPTVNCOP 120
 DB 312 AGTVYIEFTARETNCSTQNTLTADCTCKLHGLSLNCANVMYRPNKVVPTVRCQA 371
 QY 121 LGM 123
 DB 372 LDM 374
 RESULT 9
 ID_KNT1_RAT STANDARD; PRT; 430 AA.
 AC P01038; P04081;
 DT 01-NOV-1986 (Rel. 03, Created)
 DT 01-NOV-1988 (Rel. 09, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE T-kininogen I precursor (Major acute phase protein) (Alpha-1-MAP)
 DE (Thiostatin) (Contains: T-kinin).
 GN MAP1.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=86008264; PubMed=2413018;
 RA Furuto-Kato S., Matsumoto A., Kitamura N., Nakanishi S.;
 RT "Primary structures of the mRNAs encoding the rat precursors for
 RT bradykinin and T-kinin. Structural relationship of kininogens with
 RT major acute phase protein and alpha 1-cysteine proteinase
 RT inhibitor.";
 RL J. Biol. Chem. 260:12054-12059(1985).
 RN [2]
 RP SEQUENCE OF 5-430 FROM N.A., AND PARTIAL SEQUENCE.
 RX MEDLINE=86008266; PubMed=2413019;
 RA Anderson K.P., Heath E.C.;
 RT "The relationship between rat major acute phase protein and the
 RT kininogens.";

RL J. Biol. Chem. 260:12065-12071(1985).
 [3]
 RN SEQUENCE OF 7-430 FROM N.A.
 RP MEDLINE=85127561; PubMed=2578992;
 RX Cole T., Inglis A.S., Roxburgh C.M., Howlett G.J., Schreiber G.;
 RA "Major acute phase alpha 1-protein of the rat is homologous to bovine
 RT kininogen and contains the sequence for bradykinin: its synthesis is
 RT regulated at the mRNA level.";
 RL FEBS Lett. 182:57-61(1985).
 [4]
 RN SEQUENCE OF 1-65 FROM N.A.
 RP MEDLINE=87250580; PubMed=2439509;
 RX Fung W.-P., Schreiber G.;
 RA "Structure and expression of the genes for major acute phase alpha 1-
 RT protein (thiostatin) and kininogen in the rat.";
 RL J. Biol. Chem. 262:9298-9308(1987).
 CC -!- FUNCTION: Kininogens are plasma glycoproteins with a number of
 CC functions: (1) as precursor of the active peptide bradykinin and
 CC effect smooth muscle contraction, induction of hypotension and
 CC increase of vascular permeability. (2) They play a role in blood
 CC coagulation by helping to position optimally prekallikrein and
 CC factor XI next to factor XII. (3) They are inhibitor of thiol
 CC proteases.
 CC -!- SUBCELLULAR LOCATION: Extracellular.
 CC -!- TISSUE SPECIFICITY: Plasma.
 CC -!- INDUCTION: In response to an inflammatory stimulant. T-kininogen
 CC II synthesis is induced and the plasma concentration of
 CC T-kininogen I is raised.
 CC -!- PTM: AS T-KININ IS PRECEDED BY A MET INSTEAD OF AN ARG OR LYS, IT
 CC IS NOT RELEASED FROM ITS PRECURSOR BY EITHER TISSUE OR PLASMA
 CC KALLIKREIN.
 CC -!- MISCELLANEOUS: Rats express four types of kininogens: the
 CC classical HMW and LMW kininogens produced by alternative splicing
 CC of the same gene, and two additional LMW-like kininogens: T-I and
 CC T-II.
 CC -!- SIMILARITY: Contains 3 cystatin-like domains.
 CC -!- CAUTION: In addition to the conflicts described in the feature
 CC table, Ref.2 sequence differs from that shown in positions 257,
 CC 262, 268, 269, 314, 315, 331, 332 and 389. In all those
 CC positions the alternate amino acid is the one present in T-II
 CC kininogen.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
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 CC the European Bioinformatics Institute. There are no restrictions on its
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 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; M11883; AAA41489.1; -.
 CC EMBL; M11661; AAA41570.1; -.
 CC EMBL; M16454; AAA41568.1; -.
 CC EMBL; X02299; CAA26162.1; ALT_SEQ.
 CC PIR; A01286; KGRIT1.
 CC PIR; A23897; A23897.
 CC PIR; A27115; A27115.
 CC GlycoSuiteDB; P01048; -.
 CC InterPro; IPR000010; Cystatin.
 CC Pfam; PF00031; cystatin; 3.
 CC SMART; SM00043; CY; 3.
 CC PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;
 KW Thiol protease inhibitor; Bradykinin; Acute phase; Signal.
 FT SIGNAL 1 18
 FT CHAIN 19 430 KININOGEN, T-I.
 FT CHAIN 19 375 HEAVY CHAIN.
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 FT CHAIN 387 430 LIGHT CHAIN.
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FT DISULFID 83 94 BY SIMILARITY.
FT DISULFID 107 125 BY SIMILARITY.
FT DISULFID 141 144 BY SIMILARITY.
FT DISULFID 205 217 BY SIMILARITY.
FT DISULFID 228 247 BY SIMILARITY.
FT DISULFID 263 266 BY SIMILARITY.
FT DISULFID 327 339 BY SIMILARITY.
FT DISULFID 350 369 BY SIMILARITY.
FT CARBOHYD 82 92 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 126 126 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 326 326 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CONFLICT 26 28 LNC -> MDR (IN REF. 2).
FT CONFLICT 55 55 V -> L (IN REF. 2).
FT CONFLICT 61 61 E -> Y (IN REF. 1).
FT CONFLICT 83 83 C -> K (IN REF. 3).
FT CONFLICT 166 166 S -> F (IN REF. 2 AND 3).
FT CONFLICT 179 181 REV -> TKI (IN REF. 2).
FT CONFLICT 193 193 N -> D (IN REF. 2).
FT CONFLICT 212 212 S -> F (IN REF. 2).
FT CONFLICT 214 214 R -> H (IN REF. 3).
FT CONFLICT 229 229 T -> R (IN REF. 2).
FT CONFLICT 233 233 H -> Y (IN REF. 2).
FT CONFLICT 257 257 E -> S (IN REF. 2).
FT CONFLICT 262 262 R -> K (IN REF. 2).
FT CONFLICT 264 264 R -> F (IN REF. 2).
FT CONFLICT 268 268 RE -> KN (IN REF. 2).
FT CONFLICT 295 295 I -> L (IN REF. 2).
FT CONFLICT 314 315 VI -> TK (IN REF. 2).
FT CONFLICT 331 332 SK -> TN (IN REF. 2).
FT CONFLICT 389 389 R -> Q (IN REF. 2).
FT CONFLICT 414 414 R -> G (IN REF. 2 AND 3).
FT CONFLICT 415 415 A -> L (IN REF. 2).
FT CONFLICT 420 421 DH -> ER (IN REF. 3).
FT CONFLICT 430 430 P -> S (IN REF. 1).
SQ SEQUENCE 430 AA; 47715 MW; FAEBB78FAF4723C3 CRC64;

Query Match 60.8%; Score 401; DB 1; Length 430;
Best Local Similarity 61.8%; Pred. No. 6.3e-31;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGCPRDIPNTPSPELEETLTHITKLNNAENATFYFKIDNVKARVQV 60
Db 252 GDDLPELPKNCRCGPRIEVDPSPELKALGSLAQLNAQNHIFYFKIDTVKATSQV 311
QY 61 AGKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVVVPWEKKIYPTVNCQP 120
Db 312 AGVIVVIEFIARETNCQSKSTELTADCTKHLGQSLNCNANVYMRPWENKVVPTVRCQA 371
QY 121 LQM 123
Db 372 LDM 374

RESULT 10
CYTF MOUSE
ID CYTF MOUSE STANDARD; PRT; 144 AA.
AC O89098;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Cystatin F precursor (Leukocystatin) (Cystatin 7) (Cystatin-like
DE metastasis-associated protein) (CMAP).
GN CS77.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Euthera; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98298157; PubMed=9632704;
RA Halfon S., Ford J., Foster J., Dowling L., Lucian L., Sterling M.,

Xu Y., Weiss M., Ikeda M., Liggett D., Helms A., Caux C., Lebecque S.,
Haum C., Menon S., McClanahan T., Gorman D., Zurawski G.;
"Leukocystatin, a new class II cystatin expressed selectively by
hematopoietic cells.";
J. Biol. Chem. 273:16400-16408(1998).
-!- FUNCTION: Inhibits papain and cathepsin L but with affinities
lower than other cystatins. May play a role in immune regulation
through inhibition of a unique target in the hematopoietic system.
-!- SUBCELLULAR LOCATION: Secreted (Probable).
-!- SIMILARITY: Belongs to the cystatin family.
-----
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-----
EMBL; AF031826; AAC40140.1; -.
EMBL; AF031825; AAC40139.1; -.
DR HSPP; P01034; I096.
DR MGD; MGI:1298217; Cst7.
DR InterPro; IPR00010; Cystatin.
DR Pfam; PFO0031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00287; CYSTATIN; FALSE NEG.
KW Thiol protease inhibitor; Glycoprotein; Signal.
FT SIGNAL 1 18 POTENTIAL.
FT CHAIN 19 144 CYSTATIN F.
FT ACT_SITE 36 36 REACTIVE SITE.
FT SITE 80 84 SECONDARY AREA OF CONTACT.
FT DISULFID 98 109 BY SIMILARITY.
FT DISULFID 123 143 BY SIMILARITY.
SQ SEQUENCE 144 AA; 16380 MW; B5937334C1B4A89C CRC64;

Query Match 26.5%; Score 175; DB 1; Length 144;
Best Local Similarity 34.4%; Pred. No. 6.8e-10;
Matches 42; Conservative 23; Mismatches 45; Indels 8; Gaps 4;

QY 2 KDFVQPTKICVGCPRDIPNTPSPELEETLTHITKLNNAENATFYFKIDNVKARVQV 61
Db 27 KDLI---SSVKGFPTETNPGVLKARHSEVENFNCNTDIFLKESHVSKALVQVVK 83
QY 62 GKXYFIDFVARETTCSKESNEELTESC---TKLGQSLDCNAEVVVPWEKKI-YPTVN 117
Db 84 GLKYLEVIGRTTCRKTHQL-DNCDPQTNPALKRTLYCYSEVMVTPWLHSFEVPVLL 142
QY 118 CQ 119
Db 143 CQ 144

RESULT 11
CYTF HUMAN
ID CYTF HUMAN STANDARD; PRT; 145 AA.
AC O76056; Q9UED4;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Cystatin F precursor (Leukocystatin) (Cystatin 7) (Cystatin-like
DE metastasis-associated protein) (CMAP).
GN CS77.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98406133; PubMed=9733783;
RA Ni J., Fernandez M.A., Danielsson L., Chillakuru R.A., Zhang J.,
Grubb A., Su J., Gentz R., Abrahamson M.;
"Leukocystatin F is a glycosylated human low molecular weight cysteine

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RT RT proteinase inhibitor.";
RL J. Biol. Chem. 273:24797-24804 (1998).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=98298157; PubMed=9632704;
RA Halfon S., Ford J., Foster J., Dowling L., Lucian L., Sterling M.,
RA Xu Y., Weiss M., Ikeda M., Liggett D., Helms A., Caux C., Lebecque S.,
RA Hannum C., Menon S., McClanahan T., Gorman D., Zurawski G.;
RA "Leukocystatin, a new class II cystatin expressed selectively by
RT hematopoietic cells.";
RL J. Biol. Chem. 273:16400-16408 (1998).
RN [3]
RP SEQUENCE FROM N.A.
RA Morita M., Arakawa H., Yoshiuchi N.;
RT "Human homologue of murine CMAP.";
RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A.
RX MEDLINE=20399571; PubMed=10945474;
RA Morita M., Hara Y., Tanai Y., Arakawa H., Nishimura S.;
RT "Genomic construct and mapping of the gene for CMAP
RT (Leukocystatin/Cystatin F, CSTF) and identification of a proximal
RT novel gene, BSCV (C20orf3).";
RL Genomics 67:87-91 (2000).
RN [5]
RP SEQUENCE FROM N.A.
RX MEDLINE=21638749; PubMed=11780052;
RA Deloukas P., Matthews L.H., Ashurst J., Burton J., Gilbert J.G.R.,
RA Jones M., Stavrides G., Almeida J.P., Babbage A.K., Baggeley C.L.,
RA Bailey J., Barlow K.F., Bates K.N., Bead L.M., Beare D.M.,
RA Beasley O.P., Bird C.P., Blakey S.E., Bridgman A.M., Brown A.J.,
RA Buck D., Burrill W.D., Butler A.P., Carder C., Carter N.P.,
RA Chapman J.C., Clamp M., Clark G., Clark L.N., Clark S.V., Clee C.M.,
RA Clegg S., Cobley V.E., Collier R.E., Connor R.E., Corby N.R.,
RA Coulson A., Coville G.J., Deadman R., Dhami P.D., Dunn M.,
RA Ellington A.G., Frankland J.A., Fraser A., French L., Garner P.,
RA Grafham D.V., Griffiths C., Griffiths M.N.D., Gwilliam R., Hall R.E.,
RA Hammond S., Harley J.L., Heath P.D., Ho S., Holden J.L., Howden P.J.,
RA Huckle E., Hunt A.R., Hunt S.E., Jekosch K., Johnson C.M., Johnson D.,
RA Kay M.P., Kimberley A.M., King A., Knights A., Laird G.K., Lawlor S.,
RA Lehaeslalo M.H., Leversha M.A., Lloyd C., Lloyd D.M., Lovell J.D.,
RA Marsh V.L., Martin S.L., McConachie L.J., McLeay K., McMurray A.A.,
RA Milne S.A., Mistry D., Moore M.J.F., Mullikin J.C., Nickerson T.,
RA Oliver K., Parker A., Patel R., Pearce T.A.V., Peck A.I.,
RA Phillimore B.J.C.T., Prathalingam S.R., Plumb R.W., Ramsay H.,
RA Rice C.M., Ross M.T., Scott C.E., Sehra H.K., Showkneen R., Sims S.,
RA Skuce C.D., Smith M.L., Soderlund C., Steward C.A., Sulston J.E.,
RA Swann R.M., Sycamore N., Taylor R., Tee L., Thomas D.W., Thorpe A.,
RA Tracey A., Tromans A.C., Vaudin M., Wall M., Wallis J.M.,
RA Whitehead S.L., Whittaker P., Willey D.L., Williams L., Williams S.A.,
RA Wilming L., Wray P.W., Hubbard T., Durbin R.M., Bentley D.R., Beck S.,
RA Rogers J.;
RT "The DNA sequence and comparative analysis of human chromosome 20.";
RL Nature 414:865-871 (2001).
RN [6]
RP SEQUENCE FROM N.A.
RC TISSUE=Lung;
RX MEDLINE=22388257; PubMed=12477932;
RA Klausner R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Strausberg R.L., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano P.A., Peters G.J., Abramson R.D., Mullaly S.J.,
RA Boek S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettner M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,

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RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
CC -1- FUNCTION: Inhibits papain and cathepsin L but with affinities
CC lower than other cystatins. May play a role in immune regulation
CC through inhibition of a unique target in the hematopoietic system.
CC -1- SUBCELLULAR LOCATION: Secreted (Probable).
CC -1- TISSUE SPECIFICITY: Primarily expressed in peripheral blood cells
CC and spleen.
CC -1- SIMILARITY: Belongs to the cystatin family.
CC
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CC
DR EMBL; AF036342; AAC35747.1; -.
DR EMBL; AF031824; AAC39788.1; -.
DR EMBL; AB015225; BAA34941.1; ALT_INIT.
DR EMBL; AB029636; BAB11886.1; ALT_INIT.
DR EMBL; AL035661; CAB75498.1; -.
DR EMBL; BC015507; AAI15507.1; ALT_INIT.
DR HSP; F01034; IG96.
DR MIM; 603253; -.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; TAS.
DR GO; GO:0006955; P:immune response; TAS.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; Cy; 1.
DR PROSITE; PS00287; CYSTATIN; FALSE_NEG.
DR Thiol protease inhibitor; Glycoprotein; Signal.
FT SIGNAL 1 19 POTENTIAL.
FT CHAIN 20 145 CYSTATIN F.
FT ACT_SITE 37 37 REACTIVE SITE.
FT SITE 81 85 SECONDARY AREA OF CONTACT.
FT DISULFID 99 110 BY SIMILARITY.
FT DISULFID 124 144 BY SIMILARITY.
FT CARBOHYD 62 62 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 115 115 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 145 AA; 16454 MW; B2BCC4F76857CB0F CRC64;
Query Match 25.1%; Score 165.5; DB 1; Length 145;
Best Local Similarity 32.5%; Pred. No. 5.5e-09;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;
QY 9 TKTCVCPDRDPTNSPELEBTLTHITKLNAENNAFFKIDNVKARVQVWAGKYFID 68
DB 32 SRVKGPFKTIKNDPGLQAAARYSVEKFNCTNDMFLFKESRITLALQIVKGLKYLE 91
QY 69 FVARETTCSKESNEELTESCE---TKKLGQSLDCNAEVVVPWEKKI-YPTVNC 118
DB 92 VEIGRTTCKKNQHLRL-DDCDFQTNHTLKTLSCYSEVWVWPLQHFVEFVLR 144
RESULT 12
CYTM HUMAN
ID CYTM HUMAN STANDARD; PRT; 149 AA.
AC Q15828;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Cystatin M precursor (Cystatin E).
GN CST6.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;

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RN SEQUENCE FROM N.A.
 RP MEDLINE=97150844; PubMed=8995380;
 RX Sotiropoulos G., Anisowicz A., Sager R.;
 RA "Identification, cloning, and characterization of cystatin M, a novel
 RT cysteine proteinase inhibitor, down-regulated in breast cancer.";
 RL J. Biol. Chem. 272:903-910(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RP MEDLINE=97256812; PubMed=9099741;
 RA Ni J., Abrahamson M., Zhang M., Fernandez M.A., Grubb A., Su J.,
 RA Yu G.L., Li Y., Parmelee D., Xing L., Coleman T.A., Gentz S.,
 RA Thokakura R., Nguyen N., Hesselberg M., Gentz R.;
 RT "Cystatin E is a novel human cysteine proteinase inhibitor with
 RT structural resemblance to family 2 cystatins.";
 RL J. Biol. Chem. 272:10853-10858(1997).
 RN [3]
 RP SEQUENCE FROM N.A.
 RP TISSUE=Prostate;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Klausner R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Altshuler S.F., Zeberg B., Bueow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diachenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldi M.P., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahey J., Helton E., Kettelman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzyzinski M.I., Skalska U., Smalhus D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [4]
 RP CHARACTERIZATION, AND TISSUE SPECIFICITY.
 RX MEDLINE=21246880; PubMed=11348457;
 RA Zeeuwen P.L., Van Vlijmen-Willems I.M., Jansen B.J., Sotiropoulos G.,
 RA Curfs J.H., Meis J.F., Janssen J.J., Van Ruisven F., Schalkwijk J.;
 RT "Cystatin M/S expression is restricted to differentiated epidermal
 RT keratinocytes and sweat glands: a new skin-specific proteinase
 RT inhibitor that is a target for cross-linking by transglutaminase.";
 RL J. Invest. Dermatol. 116:693-701(2001).
 CC -!- FUNCTION: Shows moderate inhibition of cathepsin B but is not
 CC active against cathepsin C.
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- TISSUE SPECIFICITY: Restricted to the stratum granulosum of normal
 CC skin, the stratum granulosum/spinosum of psoriatic skin, and the
 CC secretory coils of eccrine sweat glands. Low expression levels are
 CC found in the nasal cavity.
 CC -!- PTM: Substrate for transglutaminases. Acts as an acyl acceptor but
 CC not as an acyl donor.
 CC -!- SIMILARITY: Belongs to the cystatin family.
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 CC
 DR EMBL; U62800; AAB06566.1; -.
 DR EMBL; U81233; AAB61305.1; -.
 DR EMBL; BC031334; AAB13334.1; -.
 DR HSSP; P01038; 1CEW.
 DR Genew; HGNC:2478; C5N6.

DR MIM; 601891; -.
 DR GO; GO:0004869; F:cysteine protease inhibitor activity; TAS.
 DR GO; GO:0007345; P:embryogenesis and morphogenesis; TAS.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; Cystatin; 1.
 DR SMART; SM00043; Cy1.
 DR PROSITE; PS00287; Cystatin; 1.
 KW Thiol protease inhibitor; Signal; Glycoprotein.
 FT SIGNAL 1 28 PROBALE
 FT CHAIN 29 149 CYSTATIN M.
 FT ACT_SITE 36 36 REACTIVE SITE.
 FT SITE 80 84 SECONDARY SITE OF CONTACT.
 FT DISULFID 98 113 BY SIMILARITY.
 FT DISULFID 126 146 BY SIMILARITY.
 FT CARBOHYD 137 137 N-LINKED (GLCNAC...) (POTENTIAL).
 SQ SEQUENCE 149 AA; 16511 MW; 2076A78BFC9FAC8C CRC64;
 Query Match 21.0%; Score 138.5; DB 1; Length 149;
 Best Local Similarity 31.5%; Pred. No. 21e-06;
 Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
 Qy 8 PTKICVCPDRIPDTPNSPELEETLTHITIKLNENNAFYFKIDNVKARVQVWAGKYPY 67
 Db 30 PQSRVWGLRDLSPDPQVQKAAQAAVAVSYNMGNSIYYFRDTHIIKAQSLVAGIKYPL 89
 Qy 68 DFVARETTCSKE---SNEELTESCETKLGQ--SLDCNAEVVVPWE 109
 Db 90 TMENGSTDCRKTVTGCHVDLI--TCPLAAGAQQKLRCDFFVLVVPWQ 136
 RESULT 13
 CYTC_BOVIN STANDARD; PRT; 148 AA.
 ID CYTC_BOVIN
 AC P01035; 1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 15-JUL-1999 (Rel. 38, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Cystatin C precursor (Colostrum thiol proteinase inhibitor).
 GN CST3.
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 CX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.; SEQUENCE OF 66-83, AND CHARACTERIZATION.
 RC TISSUE=Cerebrospinal fluid, and Choroid plexus;
 RX MEDLINE=98094199; PubMed=9434110;
 RA Olsson S.-L., Ek B., Wilm M., Broberg S., Rask L., Björck I.;
 RT "Molecular cloning and N-terminal analysis of bovine cystatin C
 RT identification of a full-length N-terminal region.";
 RL Biochim. Biophys. Acta 1343:203-210(1997).
 RN [2]
 RP SEQUENCE OF 37-148
 RX MEDLINE=8521205; PubMed=3891407;
 RA Hirado M., Tsunawawa S., Sakiyama F., Niinobe M., Fujii S.;
 RT "Complete amino acid sequence of bovine colostrum low-Mr cysteine
 RT proteinase inhibitor.";
 RL FEBS Lett. 186:41-45(1985).
 CC -!- FUNCTION: This is a thiol proteinase inhibitor.
 CC -!- MASS SPECTROMETRY: MW=13420; METHOD=VALDI.
 CC -!- SIMILARITY: Belongs to the cystatin family.
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 CC
 DR EMBL; Y10811; CAA71771.1; -.
 DR HSSP; P01034; 1C96.

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DR InterPro; IPR000010; Cystatin.  
DR Pfam; PF000031; cystatin; 1.  
DR SMART; SM00043; CY; 1.  
DR PROSITE; PS00287; CYSTATIN; 1.  
KW Thiol protease inhibitor; Signal; Pyroliidone carboxylic acid.  
FT SIGNAL 1 30 PROBABLE.  
FT CHAIN 31 148 CYSTATIN C.  
FT MOD_RES 31 31 PYROLIDONE CARBOXYLIC ACID (PROBABLE).  
FT ACT_SITE 40 40 REACTIVE SITE.  
FT SITE 84 88 SECONDARY AREA OF CONTACT.  
FT DISULFID 102 112 BY SIMILARITY.  
FT DISULFID 126 146 BY SIMILARITY.  
SQ SEQUENCE 148 AA; 16265 MW; EE740FE37CFB9F0E CRC64;  
  
Query Match 20.6%; Score 136; DB 1; Length 148;  
Best Local Similarity 30.6%; Pred. No. 3.7e-06;  
Matches 34; Conservative 22; Mismatches 31; Indels 24; Gaps 5;  
  
QY 22 NSPELEETLTHITKLNAENNAFFKIDNVKARQVAVGKVFIDFVARETTCSKESN 81  
DB 48 NEEGVOEALSPAVSEFNKRSDAYOSRVVRVRAKQVVGMYFLDVELGRITCTK--S 105  
QY 82 BELTESC-----ETKLGSLDCNAEVVVPWEKKIYPTVN-----CQ 119  
DB 106 QANLDCSFHNPPLKREKL-----CSFQVYVPMN-----TINLVKSCQ 147  
  
RESULT 14  
CYTC MACMU  
ID CYTC MACMU STANDARD; PRT; 146 AA.  
AC O19032;  
DT 15-JUL-1998 (Rel. 36, Created)  
DT 15-JUL-1998 (Rel. 36, Last sequence update)  
DT 28-FEB-2003 (Rel. 41, Last annotation update)  
DE Cystatin C precursor.  
GN CST3.  
OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopitheciidae;  
OC Cercopitheciinae; Macaca.  
OX NCBI_TaxID=9544;  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=97054523; PubMed=8998820;  
RA Wei L.H., Walker L.C., Levy E.;  
RT "Cystatin C. Icelandic-like mutation in an animal model of  
cerebrovascular beta-amyloidosis.";  
RL Stroke 27:2080-2085(1996).  
CC -!- FUNCTION: As an inhibitor of cysteine proteinases, this protein is  
thought to serve an important physiological role as a local  
regulator of this enzyme activity.  
CC -!- SIMILARITY: Belongs to the cystatin family.  
CC  
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or send an email to license@isb-sib.ch).  
CC  
CC EMBL; U52028; AAB64051.1; -.  
DR HSP; P01034; I936.  
DR InterPro; IPR000010; Cystatin.  
DR Pfam; PF00031; cystatin; 1.  
DR SMART; SM00043; CY; 1.  
DR PROSITE; PS00287; CYSTATIN; 1.  
KW Thiol protease inhibitor; Amyloid; Signal.  
FT SIGNAL 1 26 BY SIMILARITY.  
FT CHAIN 27 146 CYSTATIN C.  
FT ACT_SITE 37 37 REACTIVE SITE.  
FT SITE 81 85 SECONDARY AREA OF CONTACT.  
FT DISULFID 99 109 BY SIMILARITY.  
FT DISULFID 123 143 BY SIMILARITY.  
SQ SEQUENCE 146 AA; 15946 MW; 08196353C0306AA3 CRC64;  
  
Query Match 20.1%; Score 132.5; DB 1; Length 146;  
Best Local Similarity 29.0%; Pred. No. 7.8e-06;  
Matches 29; Conservative 21; Mismatches 45; Indels 5; Gaps 2;  
  
QY 13 VGCPRDPTNSPELEETLTHITKLNAENNAFFKIDNVKARQVAVGKVFIDFVAR 72  
DB 36 LGGEMDASVEEGVRALDPAVSEYNKASNDVMYHSRALQVWRARKQIVAGVNY 95  
QY 73 ETTCSKESNEELTESC-----ETKLGSLDCNAEVVVPWE 109
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Db 96 RTTCTK--NQPNLDNCPFFHEQPHLKXKAFCSFQIYSVPWQ 133

Search completed: September 24, 2004, 14:09:12
Job time : 23.364 secs

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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:05:18 ; Search time 35.424 Seconds
(without alignments)
1095.549 Million cell updates/sec

Title: US-10-661-784-1
Perfect score: 660
Sequence: 1 GKDFVQPTKICVGCPRDIP.....YVVPWEKKIYFTVNCQPLGM 123

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 3155:8202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SPTREMBL 25:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_mhc:*
- 8: sp_organelle:*
- 9: sp_phase:*
- 10: sp_plant:*
- 11: sp_podent:*
- 12: sp_virus:*
- 13: sp_vertebrate:*
- 14: sp_unclassified:*
- 15: sp_virus:*
- 16: sp_bacteriap:*
- 17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	404	61.2	140	Q7YRP6	Q7YRP6 sus scrofa
2	402	60.9	423	P70517	P70517 rattus norv
3	399	60.5	430	Q63581	Q63581 rattus norv
4	175	26.5	167	Q9QW15	Q9QW15 mus musculu
5	165.5	25.1	167	Q7Z4J8	Q7Z4J8 homo sapien
6	152.5	23.1	462	Q7Z191	Q7Z191 xenopus lae
7	152.5	23.1	462	Q7SVH2	Q7SVH2 xenopus lae
8	152.5	23.1	465	Q801B5	Q801B5 xenopus lae
9	129.5	19.6	140	Q9EPX9	Q9EPX9 mus musculu
10	122.5	18.6	455	Q800S8	Q800S8 brachydanio
11	118	17.9	464	Q801Z5	Q801Z5 cyprinus ca
12	115.5	17.5	388	Q8CB17	Q8CB17 mus musculu
13	113.5	17.2	148	Q9NH95	Q9NH95 litomosoid
14	113	17.1	140	Q80Y72	Q80Y72 mus musculu
15	111	16.8	146	Q8K397	Q8K397 mus musculu
16	111	16.8	149	Q9D1B1	Q9D1B1 mus musculu

ALIGNMENTS

RESULT 1

Q7YRP6 PRELIMINARY; PRT; 140 AA.

AC Q7YRP6; 01-OCT-2003 (TrEMBLrel. 25, Created)

DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)

DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)

DE Low molecular weight kininogen (Fragment).

GN KNG.

OS Sus scrofa (Pig).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.

OX NCBI_TaxID=9823;

RN [1]

RP SEQUENCE FROM N.A.

RA Vonnahme K.A., Fernando S.C., Ross J.A., Ashworth M.D., DeSilva U., Malayer J.R., Geisert R.D.;

RA "Porcine Endometrial and Conceptus Expression of Kininogens and Plasma Kallikrein in Cyclic and Pregnant Gilts."

RT Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.

RL EMBL: AY211363; AAP85260.1; -

DR NON_TER 1

FT NON_TER 140

SQ SEQUENCE 140 AA; 15650 MW; 177837836603F777 CRC64;

Query Match 61.2%; Score 404; DB 6; Length 140;
Best Local Similarity 78.2%; Pred. No. 5.4e-33;
Matches 79; Conservative 5; Mismatches 17; Indels 0; Gaps 0;

Qy 22 NSPELETHTHTITKLNAENNATFYFKIDNVKARVQVVGKKYFDVFARETTCSKSN 81

Db 1 DSPDLEPLNHSIAKLNAENNAVFYFKIGPVEKATVQVVGKKYSIVFTARETTCSKSN 60

Qy 82 EELTSCETCKKGOSLDCAEYVYVVPWEKKIYFTVNCQPLG 122

Db 61 EELTSCETCKKPGQILKCNASVYVVPWEKKIYFTVNCQPLG 101

RESULT 2

NCBI_TaxID=10116;
 (1)
 RN SEQUENCE FROM N.A.
 RP MEDLINE=90034172; PubMed=2806908;
 RX Anderson K.P., Croyle M.L., Lingrel J.B.;
 RA "Primary structure of a gene encoding rat T-kininogen.";
 RT Gene 81:119-128(1989)
 RL
 DR EMBL; M29090; AAA42251.1; -
 DR EMBL; M29083; AAA42251.1; JOINED.
 DR EMBL; M29084; AAA42251.1; JOINED.
 DR EMBL; M29091; AAA42251.1; JOINED.
 DR EMBL; M29085; AAA42251.1; JOINED.
 DR EMBL; M29086; AAA42251.1; JOINED.
 DR EMBL; M29087; AAA42251.1; JOINED.
 DR EMBL; M29088; AAA42251.1; JOINED.
 DR EMBL; M29089; AAA42251.1; JOINED.
 DR PIR; S68034; S68034.
 DR PIR; S68035; S68035.
 DR GO; GO:0004869; F:cytosteine protease inhibitor activity; IEA.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; cystatin; 3.
 DR SMART; SM00043; Cy; 3.
 DR PROSITE; PS00287; CYSTATIN; 2.
 SQ PROSITE; PS00287; CYSTATIN; 2.
 SQ SEQUENCE 430 AA; 47618 MW; 45508DEF4BDC978C CRC64;
 Query Match 60.5%; Score 399; DB 11; Length 430;
 Best Local Similarity 61.8%; Pred. No. 6.3e-32;
 Matches 76; Conservative 13; Mismatches 34; Indels 0; Gaps 0;
 QY 1 GKDFVQPTKICVGCPRDIPITNSPELEETLTITITKLNAENNAIFYFKIDNVKARVVV 60
 DB 252 GDDLFSLLPNKRCGPRIEIPVDSPELKEALGHSIAQLNAQHNHIFYFKIDNVKATSQV 311
 QY 61 AGKYFYDFVARETTCCKSNBELTSCETKKLGSLDCAEVVVPWEKKIYPTVACQP 120
 DB 312 AGVIYVIEFTARETNCKSKGKTELTDACETKLGSLNCNANVYRPNWKNVPTVACQA 371
 QY 121 LGM 123
 DB 372 LDM 374
 RESULT 4
 Q9QWL5 PRELIMINARY; PRT; 167 AA.
 ID Q9QWL5
 AC Q9QWL5;
 DT 01-WAY-2000 (TrEMBLrel. 13, Created)
 DT 01-WAY-2000 (TrEMBLrel. 13, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE Murine CMAP (CYSTATIN F) (LEUKOCYSTATIN).
 GN MURINE CMAP OR CST7.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 (1)
 RN SEQUENCE FROM N.A.
 RP Morita M., Arakawa H., Yoshiuchi N.;
 RA "A novel cystatin-like metastasis associated gene.";
 RT Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
 RL
 RN SEQUENCE FROM N.A.
 RP STRAIN=C57BL/6J; TISSUE=Embryo;
 RC MEDLINE=21085660; PubMed=11217851;
 RX Kawai J., Shiragawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
 RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
 RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
 RA Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
 RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
 RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
 RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
 RA Schriml L.M., Straubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
 RA Sakai K., Okido T., Furuno M., Aon H., Baldarelli R., Barsi G.,

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RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,
RA Hayashizaki Y.,
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690(2001).
DR EMBL; AB015224; BAA34940.1; -.
DR EMBL; AK004420; BAB23298.1; -.
DR HSSP; P01034; 1G96.
DR MGD; MGI:1298217; Cst7.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; Cy; 1.
SQ SEQUENCE 167 AA; 18847 MW; 61F776D8445095FE CRC64;

Query Match 26.5%; Score 175; DB 11; Length 167;
Best Local Similarity 34.4%; Pred. No. 8.3e-10;
Matches 42; Conservative 23; Mismatches 49; Indels 8; Gaps 4;

Qy 2 KDFVQPTKICVGPRIPTNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWA 61
Db 50 KDLI---SSVKPGFPKTIETNPGVLKAARSHSVKFNNTDIFLFKESHVSKALVQVVK 106

Qy 62 GKYPIDFVARETTCSKESNEELTESC---TKLGSGLDCNAEVYVVPWEKTI-YPTVN 117
Db 107 GLKTNLEVKIGRTTCRKTMHQQL-DNCDFTQNPALKRTLYCYSEVWVPIWLSHFVPEVLL 165

Qy 118 CQ 119
Db 166 CQ 167

RESULT 5
Q7Z4J8 PRELIMINARY; PRT; 167 AA.
ID Q7Z4J8
AC Q7Z4J8;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin F (Leukocystatin).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Kainine N., Chen X., Rolfs A., Halleck A., Hines L., Eisenstein S.,
RA Koundinya M., Raphael J., Moreira D., Kelley T., LaBaer J., Lin Y.,
RA Phelan M., Farmer A.;
RL Submitted (AUG-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BT009825; AAP88827.1; -.
SQ SEQUENCE 167 AA; 18857 MW; E339025A5BD60177 CRC64;

Query Match 25.1%; Score 165.5; DB 4; Length 167;
Best Local Similarity 32.5%; Pred. No. 7.5e-09;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;

Qy 9 TKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWACKYFID 68
Db 54 SRVKPGFPKTIKTNDPGVQARYSVEKFNNTDMLFKESRITRALVQIVKGLKYMLE 113

Qy 69 FVARETTCSKESNEELTESC---TKLGSGLDCNAEVYVVPWEKTI-YPTVNC 118
Db 114 VEIGRTTCKKNQHLRL-DDCDFQTNHTLTKQTLSCYSEVWVVPWLQHFVPLRC 166

RESULT 6

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Q7ZV91 PRELIMINARY; PRT; 462 AA.
ID Q7ZV91
AC Q7ZV91;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Similar to fetuin B.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Embryo;
RA Klein S., Strausberg R.;
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC043891; AAH43891.1; -.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; Cy; 2.
SQ SEQUENCE 462 AA; 53185 MW; D7BAD339961739FB CRC64;

Query Match 23.1%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 5e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

Qy 8 PTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWVAGK 63
Db 142 PGVILSTCP-DCPTANEETPTITADTLIAEYKNSNNTFYFKIDHIERVRSQWVGP 200

Qy 64 KYFIDFVARETTCSKESNEELTESC 88
Db 201 SYFIQTIKETDCMKTQENVLSNC 225

RESULT 7
Q7SYH2 PRELIMINARY; PRT; 462 AA.
ID Q7SYH2
AC Q7SYH2;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin domain fetuin-like protein.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Ventral midgut;
RA Costa R.M.B., Mason J., Lee M., Amaya E., Zorn A.M.;
RA "Novel gene expression domains reveal early patterning of the Xenopus
RT endoderm.";
RL Submitted (MAR-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY260732; AAP82289.1; -.
SQ SEQUENCE 462 AA; 53186 MW; 796F92774CC27721 CRC64;

Query Match 23.1%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 5e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

Qy 8 PTKICVGCPRDPTNSPELEETLTHITKLAENNATFYFKIDNVKKARVQVWVAGK 63
Db 142 PGVILSTCP-DCPTANEETPTITADTLIAEYKNSNNTFYFKIDHIERVRSQWVGP 200

Qy 64 KYFIDFVARETTCSKESNEELTESC 88
Db 201 SYFIQTIKETDCMKTQENVLSNC 225

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RESULT 8
Q801E5 PRELIMINARY; PRT; 465 AA.
AC Q801E5;
DT 01-JUN-2003 (TREMBlrel. 24, Created)
DT 01-JUN-2003 (TREMBlrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Hypothetical histidine-rich protein (Fragment).
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22480013; PubMed=12591597;
RA Chen Y., Jurgens K., Hollemann T., Clausen M., Ramadori G.,
RA Pieler T.;
RT "Cell-autonomous and signal-dependent expression of liver and
RT intestine marker genes in pluripotent precursor cells from Xenopus
RT embryos."; 120:277-288 (2003).
RL Mech. Dev. 120:277-288 (2003).
DR EMBL; AY188284; AAC31610.1; -.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; CY; 2.
DR KW Hypothetical protein.
FT NON_TER
SQ SEQUENCE 465 AA; 53528 MW; 0B403AB4F78BFD4 CRC64;

Query Match 23.1%; Score 152.5; DB 13; Length 465;
Best Local Similarity 38.8%; Pred. No. 5e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

QY 8 PTKICVGPDPDIPNSPELEETLTHITKLNAENNATFYKIDNVKARQVQVAGK 63
DB 145 PGVILSTCP-DCPTANEETITPTITAEITLAEVKNKSNNTFYKIDHIERVRSQWVGP 203
QY 64 KYFIDFVARETTCSKESNEELTESC 88
DB 204 SFYIQTETKETDCKMTQENVLSNC 228

RESULT 9
Q9EPX9 PRELIMINARY; PRT; 140 AA.
AC Q9EPX9;
DT 01-MAR-2001 (TREMBlrel. 16, Created)
DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)
DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)
DE Cystatin C.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=BALB/c;
RA Taupin P.J., Ray J., Fischer W.H., Suhr S.T., Hakansson K., Grubb A.,
RA Gage F.H.;
RT "FGF-2-Responsive neural stem cell proliferation requires Ccrg, a novel
RT autocorine/paracrine cofactor.";
RL Neuron 28:385-397 (2000).
DR EMBL; AF311741; AAG40283.1; -.
DR HSSP; P01034; 1G96.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00287; CYSTATIN; 1.
FT CHAIN 21 140 CYSTATIN C.

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FT VARIANT 16 16 A -> G.
FT VARIANT 84 L -> P.
SQ SEQUENCE 140 AA; 15517 MW; 3A5633406DD58D785 CRC64;

Query Match 19.6%; Score 129.5; DB 11; Length 140;
Best Local Similarity 30.0%; Pred. No. 2.6e-05;
Matches 30; Conservative 21; Mismatches 44; Indels 5; Gaps 3;

QY 13 VCGPRDIPDIPNSPELEETLTHITKLNAENNATFYKIDNVKARQVQVAGKVFIDFVAR 72
DB 30 LGAPEEADANEGRRAALDFAVSEYKNGSNDAYHSGRAIQVVRKQLVAGVNFIDVEMG 89
QY 73 ETTCSKESNEELTESC---ETKKLGSLDCNAEVVVPWE 109
DB 90 RTCTK-SQTNLTD-CFFHDQPHLMKALCSFQISVPMK 127

RESULT 10
Q800S8 PRELIMINARY; PRT; 455 AA.
AC Q800S8;
DT 01-JUN-2003 (TREMBlrel. 24, Created)
DT 01-JUN-2003 (TREMBlrel. 25, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Fetuin-A.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Jia F.;
RT "Danio rerio fetuin-A.";
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AZ217758; AAC64483.1; -.
DR GO; GO:0005874; C:microtubule; IEA.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR GO; GO:0005198; F:structural molecule activity; IEA.
DR GO; GO:0007018; P:microtubule-based movement; IEA.
DR InterPro; IPR002453; Beta tubulin.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00228; TUBULIN B-AUTOREG; 1.
SQ SEQUENCE 455 AA; 50627 MW; D822872926BA2ACB CRC64;

Query Match 18.6%; Score 122.5; DB 13; Length 455;
Best Local Similarity 26.4%; Pred. No. 0.00052;
Matches 29; Conservative 22; Mismatches 44; Indels 15; Gaps 3;

QY 10 KICVGPDPDIPDIPNSPELEETLTHITKLNAENNATFYKIDNVKARQV-VVAGKKYFID 68
DB 140 KKCPDCPGLPLPLHPEKALSVNALAKFNKQSNHKSFKLMEVGRISQWMPMGOSYFTQ 199
QY 69 FVARETTCSKESNEELTES-----CETKKLG-QLSDCNAEVY 104
DB 200 FAIMETNCTKKDAPQNPEACKALCGDQATYGFCKSKVGSEPEVECEIY 249

RESULT 11
Q801Z5 PRELIMINARY; PRT; 464 AA.
AC Q801Z5;
DT 01-JUN-2003 (TREMBlrel. 24, Created)
DT 01-JUN-2003 (TREMBlrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Fetuin long form.
OS Cyprinus carpio (Common carp).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Cyprinus.

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OX NCBI_TaxID=7962;
RN [1]
RP SEQUENCE FROM N.A.
RA Teai P.-L., Chang G.-D., Huang C.-J.;
RT "Purification and cloning of carp fetuin.";
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY225965; AA074862.1; -.
DR GO; GO:0005874; C:microtubule; IEA.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR GO; GO:0005198; F:structural molecule activity; IEA.
DR GO; GO:0007018; P:microtubule-based movement; IEA.
DR InterPro; IPR002453; Beta tubulin.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00228; TUBULIN B AUTOREG; 1.
SQ SEQUENCE 464 AA; 51698 MW; 7A54F71E44050895 CRC64;

Query Match 17.9%; Score 118; DB 13; Length 464;
Best Local Similarity 24.2%; Pred. No. 0.0015;
Matches 31; Conservative 27; Mismatches 50; Indels 20; Gaps 5;

QY 12 CVCGRDPTNSPELEETLTHITKLNAENNATFYFKIDNVKARVQ-VVAGKKYFIDFV 70
DB 142 CPDPCGGLPLHDPKGLSVKTLQKFNKESDHKSYPKLMVEVGRISTQWFGSGQSFQPA 201
QY 71 ARETTCSKE---SNEELTES-----CETKLG-QSLDCNAEYVVV-----PKEKK 111
DB 202 IMETNCTNKEAPQNEESKALCGEKARYGFKSTKVGIEPEVECEIYEAKNITHPWKHP 261
QY 112 IYPTVNCQ 119
DB 262 AQSRDCK 269

RESULT 12
Q8CB17 PRELIMINARY; PRT; 388 AA.
AC Q8CB17;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DR Fetuin beta.
GN FETUB.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=C57BL/6J; TISSUE=Vagina;
RX MEDLINE=22354683; PubMed=12466851;
RA The FANTOM Consortium,
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs.";
RL Nature 420:563-573(2002).
DR EMBL; AK037043; BAC29682.1; -.
DR MGD; MGI:1890221; Fetub.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR InterPro; IPR001363; Fetuin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; CY; 2.
DR PROSITE; PS01254; FETUIN 1; 1.
DR PROSITE; PS01255; FETUIN 2; 1.
SQ SEQUENCE 388 AA; 42742 MW; 78CFAD73A8D8DC22 CRC64;

Query Match 17.5%; Score 115.5; DB 11; Length 388;
Best Local Similarity 26.7%; Pred. No. 0.0022;
Matches 31; Conservative 24; Mismatches 49; Indels 13; Gaps 4;

QY 12 CVCGRDPTNSPELEETLTHITKLNAENNATFYFKIDNVKARVQVWAGKKYFIDFVA 71
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DB 154 CPDCSPIDLNSALEAATESLAKFNKSPSKY-ELUVKTKANNQVSPAYVEYLI 212
QY 72 RETTCSK-----ESNEELTESCETKLGQSLDCNAEYVVVWPKKIYPTVNCQ 119
DB 213 KEAPCTKQASCSLQHSDEPVGICQGSTVQSLL--RHVPLIQPKSV--TVTCE 264

RESULT 13
Q9NH95 PRELIMINARY; PRT; 148 AA.
AC Q9NH95;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Ls-cystatin.
OS Litomosoides sigmodontis.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
OC Ouchocercidae; Litomosoides.
OX NCBI_TaxID=42156;
RN [1]
RP SEQUENCE FROM N.A.
RA Pfaff A.W., Hoffmann W.H., Taylor D.W., Schulz-Key H.;
RT "Characterization and immunological properties of a cysteine protease
RT inhibitor of the filarial parasite Litomosoides sigmodontis.";
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF229173; AAF35896.1; -.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
DR PROSITE; PS00287; CYSTATIN; 1.
FT CHAIN 25 148 LS-CYSTATIN.
SQ SEQUENCE 148 AA; 16686 MW; 2950AA89CA5339C9 CRC64;

Query Match 17.2%; Score 113.5; DB 5; Length 148;
Best Local Similarity 34.1%; Pred. No. 0.0011;
Matches 30; Conservative 16; Mismatches 37; Indels 5; Gaps 3;

QY 25 ELEELTTHITKLNAENNATFYFKIDNVKARVQVWAGKKYFIDFVARETTCSKESNEEL 84
DB 49 EIQEMLP SILTKVNCQSDAHLMPIKVLKVSQVWAGKKYKVEIQVAESDCKKSNEKI 108
QY 85 -TESCETKLGQSLD--CNAEYVVVWPE 109
DB 109 DLKTC--KKLEGHDPQIITLEVWEKWE 134

RESULT 14
Q80Y72 PRELIMINARY; PRT; 140 AA.
AC Q80Y72;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cystatin-like 1.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA TISSUE=testicle;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M.J., Udell T.B., Toshiyuki S., Carninci P., Frange C.,
RA Brownstein M.J., Loquellano N.A., Peters G.J., Mullany S.J.,
RA Raha S.S., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Bosak S.A.,
```


GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:06:08 ; Search time 13.284 Seconds
(without alignments)
890.662 Million cell updates/sec

Title: US-10-661-784-1

Perfect score: 660

Sequence: 1 GKDFVQPTKICVGCPRDIP.....YVFWERKIYPTVNCQPLGM 123

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 293366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : PIR 78.*

1: pir1.*

2: pir2.*

3: pir3.*

4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	660	100.0	427	1 KGHUL1	kininogen, LMW pre
2	660	100.0	644	1 KGHUL1	kininogen, LMW pre
3	477	72.3	436	1 KGBOL1	kininogen, LMW I p
4	477	72.3	621	1 KGBOL1	kininogen, LMW I p
5	450	68.2	434	1 KGBOL2	kininogen, LMW II
6	450	68.2	619	1 KGBOL2	kininogen, LMW II
7	426	64.5	433	2 A28055	K-kininogen, LMW I
8	426	64.5	639	2 A25486	kininogen, LMW I p
9	409	62.0	430	2 A28997	major acute phase
10	409	62.0	430	2 B28055	T-kininogen, LMW I
11	402	60.9	423	1 KGRTM	major acute phase
12	401	60.8	430	1 KGRTT1	T-kininogen I prec
13	136	20.6	112	1 UPBO	cystatin - bovine
14	133	20.2	91	2 S68034	T-kininogen (clone
15	133	20.2	91	2 S68035	T-kininogen (clone
16	130	19.7	127	2 S07085	cystatin C precurs
17	129	19.5	120	2 S10587	cystatin C - rat
18	128	19.4	111	2 A28793	cystatin - puff ad
19	127.5	19.3	140	2 A36163	cystatin C precurs
20	127.5	19.3	146	1 UNHU	cystatin C precurs
21	125	18.9	141	2 B29632	cystatin SA precur
22	118.5	18.0	139	1 UDCB	cystatin precursor
23	113.5	17.2	122	2 A34644	sarcocystatin A pr
24	113	17.1	141	2 JG1470	cystatin S precurs
25	111.5	16.9	111	1 JG2040	cystatin - chum sa
26	109.5	16.6	141	1 UNHUP2	cystatin SN precur
27	109	16.5	141	1 UNHUP1	cystatin S precurs
28	107.5	16.3	132	2 JG4918	cystatin precursor
29	107	16.2	139	2 T33740	hypothetical prote

RESULT 1

KGHUL1

kininogen, LMW precursor [validated] - human

N;Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen

N;Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen

C;Species: Homo sapiens (man)

C;Date: 06-Jul-1982 #sequence revision 27-Nov-1985 #text change 08-Dec-2000

C;Accession: A01280; B25276; A27900; A27699; A31905; A34030

R;Okubo, I.; Kurachi, K.; Takasawa, T.; Shikawa, H.; Sasaki, M.

Biochemistry 23, 5631-5697, 1984

A;Title: Isolation of a human cDNA for alpha-2-thiol proteinase inhibitor and its identi

A;Reference number: A90490; MUID:85122621; PMID:6441591

A;Accession: A01280

A;Molecule type: mRNA

A;Residues: 1-427 <OHK>

A;Cross-references: GB:K02566; NID:gl77889; PIDN:AAA35497.1; PID:gl77890

R;Takagaki, Y.; Kitamura, N.; Nakanishi, S.

J. Biol. Chem. 260, 8601-8609, 1985

A;Title: Cloning and sequence analysis of cDNAs for human high molecular weight and low

A;Reference number: A92544; MUID:85234582; PMID:2989293

A;Accession: B25276

A;Molecule type: mRNA

A;Residues: 1-427 <TAK>

A;Cross-references: GB:M11437; NID:gl86751; PIDN:AAB59551.1; PID:g386853

R;Lottspeich, F.; Kellermann, J.; Henschen, A.; Rauth, G.; Mueller-Esterl, W.

in Kinins IV, part A, Greenbaum, L.M., and Margolius, H.S., eds., pp.91-95, Plenum, New

A;Title: Amino acid sequence of the light chain of human low molecular mass kininogen.

A;Reference number: A27900

A;Accession: A27900

A;Molecule type: protein

A;Residues: 390-427 <LOT>

R;Mindrou, T.; Carretero, O.A.; Proud, D.; Walz, D.; Scicli, A.G.

Biochem. Biophys. Res. Commun. 152, 519-526, 1988

A;Title: A new kinin moiety in human plasma kininogens.

A;Reference number: A27699; MUID:88209021; PMID:3365237

A;Accession: A27699

A;Molecule type: protein

A;Residues: 380-389 <MIN>

R;Maeda, H.; Matsumura, Y.; Kato, H.

J. Biol. Chem. 263, 16051-16054, 1988

A;Title: Purification and identification of [hydroxypropyl(3)]-lysyl-bradykinin in ascitic fluid

A;Reference number: A31905; MUID:89034061; PMID:3182782

A;Accession: A31905

A;Molecule type: protein

A;Residues: 381-389 <MAE>

R;Sasaguri, M.; Ikeda, M.; Ideishi, M.; Arakawa, K.

Biochem. Biophys. Res. Commun. 150, 511-516, 1988

A;Title: Identification of [hydroxyproline(3)]-1-lysyl-bradykinin released from human plas

A;Reference number: A34030; MUID:88106632; PMID:3337729

A;Accession: A34030

A;Molecule type: protein

A;Residues: 380-389 <SAS>

R; Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
 J. Biol. Chem. 260, 8610-8617, 1985
 A; Title: Structural organization of the human kininogen gene and a model for its evolution
 A; Reference number: A92545; MUID: 85234583; PMID: 2989294
 A; Contents: annotation, gene organization
 R; Pierce, J.V.
 Fed. Proc. 27, 52-57, 1968
 A; Title: Structural features of plasma kinins and kininogens.
 A; Reference number: A91455; MUID: 90255622; PMID: 4952632
 A; Contents: annotation, bradykinin
 C; Comment: The LMW kininogen precursor is produced from the same gene as the HMW form (see C; Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the C; Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, and xiprolone residue is present in the kininogen prior to the release of bradykinin.
 C; Genes: KNG
 A; Gene: GDB:KNG
 A; Cross-references: GDB:125256; OMIM:228960
 A; Map position: 3q27-3q27
 A; Introns: 65/3; 102/3; 131/1; 188/3; 224/3; 253/1; 310/3; 346/3; 375/3; 401/3
 C; Superfamily: kininogen; cystatin homology
 C; Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glycosylation
 F; 18/Domain: signal sequence #status predicted <SIG>
 F; 19-427/Product: LMW prokininogen (kininogen I) #status predicted <MAT>
 F; 19-389, 390-427/Product: LMW kininogen II #status predicted <MAT2>
 F; 19-379/Product: LMW kininogen heavy chain #status predicted <HCH>
 F; 19-131/Domain: cystatin homology <CY1>
 F; 142-253/Domain: cystatin homology <CY2>
 F; 264-375/Domain: cystatin homology <CY3>
 F; 380-389/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>
 F; 381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
 F; 390-427/Product: LMW kininogen light chain #status experimental <LCH>
 F; 19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status predicted
 F; 28-407, 83-94, 107-126, 142-145, 206-218, 229-248, 264-267, 328-340, 351-370/Disulfide bonds:
 F; 48, 169, 205, 294/Binding site: carbonylate (Asn) (covalent) #status predicted
 F; 379-380/Cleavage site: Met-Lys (kallikrein) #status experimental
 F; 383/Modified site: 4-hydroxyproline (Pro) (partial) #status experimental
 F; 389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
 F; 401/Binding site: carbonylate (Thr) (covalent) #status absent

Query Match 100.0%; Score 660; DB 1; Length 427;
 Best Local Similarity 100.0%; Pred. No. 3.3e-55;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVPPKICVCGPRDPTNSPELEETLTHITKLAENNAATFYKIDNVKARVQV 60
 Db 253 GKDFVPPKICVCGPRDPTNSPELEETLTHITKLAENNAATFYKIDNVKARVQV 312
 QY 61 AGKYPIDFVARETTCKESNELTESCTKLGSLDCAEYVVPWEKKIYPTVNCQP 120
 Db 313 AGKYPIDFVARETTCKESNELTESCTKLGSLDCAEYVVPWEKKIYPTVNCQP 372
 QY 121 LGM 123
 Db 373 LGM 375

RESULT 2
 KGHU1
 N; kininogen, HMW precursor [validated] - human
 N; Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen; prokininogen
 N; Contains: bradykinin (kallidin I); HMW kininogen I; HMW kininogen II; low molecular weight kininogen
 C; Species: Homo sapiens (man)
 C; Date: 28-May-1986 #sequence_revision 28-May-1986 #text_change 08-Dec-2000
 C; Accession: A01279; S32422; A91153; A24871; A27899; A31905; A34030; S02
 R; Ohkubo, I.; Kurachi, K.; Takasawa, T.; Shiohara, H.; Sasaki, M.
 Biochemistry 23, 5691-5697, 1984
 A; Title: Isolation of a human cDNA for alpha-2-thiol proteinase inhibitor and its identification
 A; Reference number: A90490; MUID: 85122621; PMID: 6441591
 A; Accession: A01279
 A; Molecule type: mRNA
 A; Residues: 1-389 <OHK>
 A; Cross-references: GB:K02566; NID: g177889
 R; Takagaki, Y.; Kitamura, N.; Nakanishi, S.

J. Biol. Chem. 260, 8601-8609, 1985
 A; Title: Cloning and sequence analysis of cDNAs for human high molecular weight and low molecular weight kininogen
 A; Reference number: A92544; MUID: 85234582; PMID: 2989293
 A; Accession: A25276
 A; Molecule type: mRNA
 A; Residues: 1-592, '1', 594-644 <TAK>
 A; Cross-references: GB:M11437; NID: g186751; PIDN: AAB59550.1; PID: g386852
 R; Auerwald, E.A.; Roessler, D.; Mentele, R.; Assfalg-Machleidt, I.
 FEBS Lett. 321, 93-97, 1993
 A; Title: Cloning, expression and characterization of human kininogen domain 3.
 A; Reference number: S32422; MUID: 93223854; PMID: 8467916
 A; Accession: S32422
 A; Molecule type: mRNA
 A; Residues: 'ANSM', 253-377 <AUE>
 A; Note: differences are due to known cloning artifacts
 R; Lottspeich, F.; Kellermann, J.; Henschen, A.; Foersts, B.; Muller-Esterl, W.
 Eur. J. Biochem. 152, 307-314, 1985
 A; Title: The amino acid sequence of the light chain of human high-molecular-mass kininogen
 A; Reference number: A91153; MUID: 86030270; PMID: 4054110
 A; Accession: A91153
 A; Molecule type: protein
 A; Residues: 379-644 <LOP>
 A; Note: the bradykinin sequence preceding the light chain sequence was not determined in R; Kellermann, J.; Lottspeich, F.; Henschen, A.; Mueller-Esterl, W.
 Eur. J. Biochem. 154, 471-478, 1986
 A; Title: Completion of the primary structure of human high-molecular-mass kininogen. The
 A; Reference number: A24871; MUID: 86108361; PMID: 3484703
 A; Accession: A24871
 A; Molecule type: protein
 A; Residues: '2', 20-380 <KEL1>
 R; Kellermann, J.; Lottspeich, F.; Henschen, A.; Mueller-Esterl, W.
 In: Kinins IV, Greenbaum, L.M., and Margolius, H.S., ed., pp. 85-89, Plenum Press, New York
 A; Title: Amino acid sequence of the light chain of human high molecular mass kininogen.
 A; Reference number: A27899
 A; Accession: A27899
 A; Molecule type: protein
 A; Residues: 379-389, 'K', 390-407, 'Q', 409-644 <KEL2>
 R; Miroslav, T.; Carrettero, O.A.; Proud, D.; Walz, D.; Scicli, A.G.
 Biochem. Biophys. Res. Commun. 152, 519-526, 1988
 A; Title: A new kinin moiety in human plasma kininogens.
 A; Reference number: A27899; MUID: 88209021; PMID: 3365237
 A; Accession: A27899
 A; Molecule type: protein
 A; Residues: 380-389 <MIN>
 R; Maeda, H.; Matsumura, Y.; Kato, H.
 J. Biol. Chem. 263, 16051-16054, 1988
 A; Title: Purification and identification of [hydroxyprolyl(3)]bradykinin in ascitic fluid
 A; Reference number: A31905; MUID: 89034061; PMID: 3182782
 A; Accession: A31905
 A; Molecule type: protein
 A; Residues: 381-389 <MAE>
 R; Sasaguri, M.; Ikeda, M.; Ideishi, M.; Arakawa, K.
 Biochem. Biophys. Res. Commun. 150, 511-516, 1988
 A; Title: Identification of [hydroxyproline(3)]-lysyl-bradykinin released from human plasma
 A; Reference number: A34030; MUID: 88106632; PMID: 3337729
 A; Accession: A34030
 A; Molecule type: protein
 A; Residues: 380-389 <SAS>
 R; Lenaric, B.; Gabrijelcic, D.; Rozman, B.; Drobnic-Kosorok, M.; Turk, V.
 Biol. Chem. Hoppe-Seyler 369, 257-261, 1988
 A; Title: Human cathepsin B and cysteine proteinase inhibitors (CPIs) in inflammatory and
 A; Reference number: S02482; MUID: 89076517; PMID: 3264507
 A; Accession: S02482
 A; Molecule type: protein
 A; Residues: 1-19; 189-192; 310-314; 381-389 <LEN1>
 R; Kato, H.; Matsumura, Y.; Maeda, H.
 FEBS Lett. 232, 252-254, 1988
 A; Title: Isolation and identification of hydroxyproline analogues of bradykinin in human
 A; Reference number: A61495; MUID: 88211869; PMID: 3366244
 A; Accession: A61495
 A; Molecule type: protein
 A; Residues: 380-389 <KAT1>
 A; Experimental source: urine

A>Note: this peptide had Pro-383 modified to 4-hydroxyproline

A:Accession: B61495
 A:Molecule type: protein
 A:Residues: 381-389 <KAT2>
 A:Experimental source: urine
 A:Note: this peptide had Pro-383 modified to 4-hydroxyproline
 A:Accession: C61495
 A:Molecule type: protein
 A:Residues: 380-389 <KAT3>
 R:Lenarcic, B.; Krasovec, M.; Ritonja, A.; Olafsson, I.; Turk, V.
 FEBS Lett. 280, 211-215, 1991
 A>Title: Inactivation of human cystatin C and kininogen by human cathepsin D.
 A:Reference number: S14303; MUID:91192133; PMID:2013314
 A:Accession: S14447
 A:Molecule type: protein
 A:Residues: 264-359, 'N', 361-375 <LEN2>
 R:Little, S.S.; Johnson, D.A.
 Biochem. J. 307, 341-346, 1995
 A>Title: Human mast cell tryptase isoforms: separation and examination of substrate-specificity
 A:Reference number: S55239; MUID:95251593; PMID:7733867
 A:Accession: S55239
 A:Molecule type: protein
 A:Residues: 450-452, 'X', 454, 'X', 456 <JIT>
 R:Straczek, J.; Maachi, F.; le Nguyen, D.; Becchi, M.; Heulin, M.H.; Nabat, P.; Bellevil
 FEBS Lett. 373, 207-211, 1995
 A>Title: Purification from human plasma of a tetrapeptide that potentiates insulin-like
 A:Reference number: S68059; MUID:96033974; PMID:7589467
 A:Accession: S68059
 A:Molecule type: protein
 A:Residues: 431-434 <STR>
 R:Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
 J. Biol. Chem. 260, 8610-8617, 1985
 A>Title: Structural organization of the human kininogen gene and a model for its evolution
 A:Reference number: A92545; MUID:85234583; PMID:2989294
 A:Contents: annotation; gene organization
 R:Pierce, J.V.
 Fed. Proc. 27, 52-57, 1968
 A>Title: Structural features of plasma kinins and kininogens.
 A:Reference number: A91455; MUID:90255622; PMID:4952632
 A:Contents: annotation; bradykinin
 C:Comment: The HMW kininogen precursor and the LMW form are produced from the same gene
 C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
 C:Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is impo
 C:Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i
 xyproline residue is present in the kininogen prior to the release of bradykinin.
 C:Genetics:
 A:Gene: GDB:KNG
 A:Cross-references: GDB:125256; OMIM:228960
 A:Map position: 3q27.3q27
 A:Introns: 65/3; 102/3; 131/1; 188/3; 224/3; 253/1; 310/3; 346/3; 375/3
 C:Superfamily: kininogen; cystatin homology
 C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupl
 F:1-18/Domain: signal sequence #status experimental <SIG>
 F:19-644/Product: HMW kininogen I (prokininogen) #status experimental <MAT1>
 F:19-379,390-644/Product: HMW kininogen II #status experimental <MAT2>
 F:19-379/Domain: HMW kininogen heavy chain #status experimental <HC>
 F:19-379/Domain: HMW kininogen heavy chain #status experimental <CY1>
 F:19-131/Domain: cystatin homology <CY2>
 F:142-253/Domain: cystatin homology <CY2>
 F:264-375/Domain: cystatin homology <CY3>
 F:380-389/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>
 F:381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
 F:390-644/Domain: HMW kininogen light chain #status experimental <LCH>
 F:421-510/Region: glycine/histidine/lysine-rich 30-residue repeats
 F:19/Modified site: low molecular weight promoting factor #status experimental <GF
 F:28-614,83-94,107-126,145-206-218,229-248,264-267,328-340,351-370/Disulfide bonds:
 F:48/Binding site: carboxylate (Asn) (covalent) #status absent
 F:169,205,294/Binding site: carboxylate (Asn) (covalent) #status experimental
 F:379-380/Cleavage site: Met-Lys (kallikrein) #status experimental
 F:383/Modified site: 4-hydroxyproline (Pro) (partial) #status experimental
 F:389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
 F:401,533,542,546,557,571,593,628/Binding site: carboxylate (Thr) (covalent) #status ex
 F:577/Binding site: carboxylate (Ser) (covalent) #status experimental

Query Match 100.0%; Score 660; DB 1; Length 644;

Best Local Similarity 100.0%; Pred. NO. 5.1e-55;
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GKDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKKARQVYV 60

Db 253 GKDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKKARQVYV 312

QY 61 AGKKYFIDFVARETTCESKESNEELTESCTKKLGSLDCNAEYVYVVPWEKKIYPTVNCQPL 120

Db 313 AGKKYFIDFVARETTCESKESNEELTESCTKKLGSLDCNAEYVYVVPWEKKIYPTVNCQPL 372

QY 121 LGM 123

Db 373 LGM 375

RESULT 3

KGBOL1

kininogen, LMW I precursor - bovine

N;Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen

N;Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen

C;Species: Bos primigenius taurus (cattle)

C;Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999

C;Accession: A01283

R;Nawa, H.; Kitamura, N.; Hirose, T.; Asai, M.; Inayama, S.; Nakanishi, S.

Proc. Natl. Acad. Sci. U.S.A. 80, 90-94, 1983

A>Title: Primary structures of bovine liver low molecular weight kininogen precursors an

A;Reference number: A93984; MUID:83117859; PMID:6572010

A;Accession: A01283

A:Molecule type: mRNA

A;Residues: 1-436 <NAW>

A;Cross-references: GB:J00010; GB:V00426; NID:G163256; PID:AAA30604.1; PID:G163257

C;Comment: The LMW kininogen precursor is produced from the same gene as the HMW form as

C;Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the

C;Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i

xyproline residue is present in the kininogen prior to the release of bradykinin.

C;Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glyc

F:1-18/Domain: signal sequence #status predicted <SIG>

F:19-436/Product: LMW kininogen I #status predicted <MAT>

F:19-378/Product: LMW kininogen I heavy chain #status predicted <HC>

F:19-130/Domain: cystatin homology <CY1>

F:141-252/Domain: cystatin homology <CY2>

F:263-374/Domain: cystatin homology <CY3>

F:379-388/Product: lysyl-bradykinin (kallidin II) #status predicted <KBDY>

F:380-388/Product: bradykinin (kallidin I) #status predicted <BDY>

F:389-436/Product: LMW kininogen I light chain #status experimental <LCH>

F:19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status predicted

F:27-406,82-93,106-125,141-144,205-217,228-247,263-286,327-339,350-369/Disulfide bonds:

F:47,87,168,169,197,204/Binding site: carboxylate (Asn) (covalent) #status predicted

F:378-379/Cleavage site: Met-Lys (kallikrein) #status predicted

F:382/Modified site: 4-hydroxyproline (Pro) #status predicted

F:388-389/Cleavage site: Arg-Ser (kallikrein) #status predicted

Query Match 72.3%; Score 477; DB 1; Length 436;

Best Local Similarity 71.9%; Pred. NO. 9.2e-38;

Matches 87; Conservative 14; Mismatches 20; Indels 0; Gaps 0;

QY 2 KDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKKARQVYV 61

Db 253 KDFVQPPTKICVGCPRDIPNTSPLEETLTHITIKLNAENNAFFYFKIDNVKKARQVYV 312

QY 62 GKXYFIDFVARETTCESKESNEELTESCTKKLGSLDCNAEYVYVVPWEKKIYPTVNCQPL 121

Db 313 GLKYSIVFARETTCESKESNEELTESCTKKLGSLDCNAEYVYVVPWEKKIYPTVNCQPL 372

QY 122 G 122

Db 373 G 373

RESULT 4

KGSOH1
kininogen, HMW I precursor - bovine
N/Alternate names: alpha-2-thiol proteinase inhibitor; prokininogen
N/Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C/Species: Bos primigenius taurus (cattle)
C/Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999
C/Accession: A01281; A91923; A91938; A29559
R/Kitamura, N.; Takagaki, Y.; Furuto, S.; Tanaka, T.; Nawa, H.; Nakanishi, S.
Nature 305, 545-549, 1983
A/Title: A single gene for bovine high molecular weight and low molecular weight kininogen
A/Reference number: A93317; MUID:84014106; PMID:6571699
A/Accession: A01281
A/Molecule type: mRNA
A/Residues: 1-621 <KIT>
A/Cross-references: GB:V01491; GB:K01757; NID:G491; PIDN:CAA24735.1; PID:G492
R/Kato, H.; Nagasawa, S.; Suzuki, T.
J. Biochem. 67, 313-323, 1970
A/Title: Studies on the structure of bovine kininogen: cleavages of disulfide bonds and
A/Reference number: A91923; MUID:70180420; PMID:4986212
A/Accession: A91923
A/Molecule type: protein
A/Residues: 378-393 <KAT>
R/Han, Y.N.; Komiya, M.; Iwanaga, S.; Suzuki, T.
J. Biochem. 77, 55-68, 1975
A/Title: Studies on the primary structure of bovine high-molecular-weight kininogen. Am
A/Reference number: A91938; MUID:75170265; PMID:1169237
A/Accession: A91938
A/Molecule type: protein
A/Residues: 458-498 <HAN>
R/Sueyoshi, T.; Miyata, T.; Hashimoto, N.; Kato, H.; Hayashida, H.; Miyata, T.; Iwanaga,
J. Biol. Chem. 262, 2768-2779, 1987
A/Title: Bovine high molecular weight kininogen. The amino acid sequence, positions of c
A/Reference number: A92627; MUID:87137530; PMID:3546295
A/Accession: A29559
A/Molecule type: protein
A/Residues: 2', 20-123, 1', 125-127, 1', 129-378 <SUE>
R/Lottspeich, F.; Kellermann, F.; Henschel, A.; Foertsch, B.; Muller-Esterl, W.
Eur. J. Biochem. 152, 307-314, 1985
A/Title: The amino acid sequence of the light chain of human high-molecular-mass kininogen
A/Reference number: A91153; MUID:86030270; PMID:4054110
A/Contents: annotation; bovine cleavage sites; bovine carbohydrate binding sites
R/Sueyoshi, T.; Miyata, T.; Kato, H.; Iwanaga, S.
Seikagaku 56, 808, 1984
A/Title: Disulfide bonds in bovine HMW kininogens.
A/Reference number: A94300
A/Contents: annotation; disulfide bonds
A/Note: article in Japanese
C/Comment: The HMW kininogen precursor is produced from the same gene as the LMW form as
C/Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is impo
C/Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i
xyproline residue is present in the kininogen prior to the release of bradykinin.
C/Superfamily: kininogen; cystatin homology
C/Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupl
F/1-18/Domain: signal sequence #status predicted <SIG>
F/19-621/Product: HMW prokininogen I #status predicted <MAT>
F/19-379/Product: HMW kininogen I heavy chain #status experimental <HCH>
F/19-130/Domain: cystatin homology <CY1>
F/141-252/Domain: cystatin homology <CY2>
F/379-388/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>
F/380-388/Product: bradykinin (kallidin I) #status experimental <BDY>
F/389-621/Product: HMW kininogen I light chain #status experimental <LCH>
F/147-488/Region: glycine/histidine/lysine-rich
F/19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status experimen
F/27-168, 169, 204/Binding site: carbohydrate (Asn) (covalent) #status experimental
F/136/Binding site: carbohydrate (Thr) (covalent) (partial) #status experimental
F/197/Binding site: carbohydrate (Asn) (covalent) (partial) #status experimental
F/378-379/Cleavage site: Met-Lys (kallikrein) #status experimental
F/382/Modified site: 4-hydroxyproline (Pro) #status predicted
F/388-389/Cleavage site: Arg-Ser (kallikrein) #status experimental

F/398, 406 512/Binding site: carbohydrate (Ser) (covalent) #status experimental
F/399, 400, 520, 524, 536, 548, 553, 570/Binding site: carbohydrate (Thr) (covalent) #status ex
F/498-499/Cleavage site: Arg-Thr (kallikrein) #status experimental

Query Match 72.3%; Score 477; DB 1; Length 621;
Best Local Similarity 71.9%; Pred. No. 1.4e-37;
Matches 87; Conservative 14; Mismatches 20; Indels 0; Gaps 0;

Qy 2 KDVQPTKICVCPDIPNPSLEETLTHITKLNANNATFFYFKIDNVKARVQVVA 61

Db 253 KDVQPTKLCACCPKIPVDSPLSEPLSHSIKLNNEHGFYFKIDTVKATVQVVA 312

Qy 62 GKXFDIFVARETTCESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTVNCQPL 121

Db 313 GLXYSIVFIARETTCESKGSNEELTKSCINIHQILHCDANVVPWEKKIYPTVNCQPL 372

Qy 122 G 122

Db 373 G 373

RESULT 5

KGBO12

kininogen, LMW II precursor - bovine

N/Alternate names: alpha-2-thiol proteinase inhibitor; prokininogen

N/Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen

C/Species: Bos primigenius taurus (cattle)

C/Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 28-May-1999

C/Accession: A01284

R/Nawa, H.; Kitamura, N.; Hirose, T.; Asai, M.; Inayama, S.; Nakanishi, S.

Proc. Natl. Acad. Sci. U.S.A. 80, 90-94, 1983

A/Title: Primary structures of bovine liver low molecular weight kininogen precursors and
A/Reference number: A93984; MUID:83117859; PMID:6572010

A/Accession: A01284

A/Molecule type: mRNA

A/Residues: 1-434 <NAW>

A/Cross-references: GB:V00427; GB:J00011; NID:G489; PIDN:CAA23710.1; PID:G490

C/Comment: The LMW kininogen precursor is produced from the same gene as the HMW form as

C/Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the

C/Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i

xyproline residue is present in the kininogen prior to the release of bradykinin.

C/Superfamily: kininogen; cystatin homology

C/Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glyco

F/1-18/Domain: signal sequence #status predicted <SIG>

F/19-434/Product: LMW kininogen I #status predicted <MAT>

F/19-377/Product: LMW kininogen I heavy chain #status predicted <HCH>

F/19-130/Domain: cystatin homology <CY1>

F/141-252/Domain: cystatin homology <CY2>

F/261-372/Domain: cystatin homology <CY3>

F/377-386/Product: lysyl-bradykinin (kallidin II) #status predicted <KBDY>

F/378-386/Product: bradykinin (kallidin I) #status predicted <BDY>

F/387-434/Product: LMW kininogen I light chain #status experimental <LCH>

F/19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status predicted

F/27-404, 82-93, 106-125, 141-144, 205-217, 228-247, 261-264, 325-337, 348-367/Disulfide bonds: #

F/47, 87, 168, 169, 197, 204, 280/Binding site: carbohydrate (Asn) (covalent) #status predicted

F/376-377/Cleavage site: Met-Lys (kallikrein) #status predicted

F/380/Modified site: 4-hydroxyproline (Pro) #status predicted

F/386-387/Cleavage site: Arg-Ser (kallikrein) #status predicted

Query Match 68.2%; Score 450; DB 1; Length 434;

Best Local Similarity 68.9%; Pred. No. 3.4e-35;

Matches 84; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

Qy 1 GKDFVQPTKICVCPDIPNPSLEETLTHITKLNANNATFFYFKIDNVKARVQV 60

Db 252 GEDFL--PMVCGCCPKPIPVDSPLSEPLSHSIKLNNEHGFYFKIDTVKATVQV 309

Qy 61 AGKFFIDFVARETTCESNEELTESCETKLGQSLDCNAEYVVPWEKKIYPTVNCQPL 120

Db 310 GGLKYSIVFIARETTCESKGSNEELTKSCINIHQILHCDANVVPWEKKIYPTVNCQPL 369

Qy 121 LG 122

Db 370 LG 371

RESULT 6

K030H2

kininogen, HMW II precursor - bovine
N:Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N:Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C:Species: Bos primigenius taurus (cattle)
C>Date: 14-Nov-1983 #sequence revision 14-Nov-1983 #text_change 22-Jun-1999
C:Accession: A01282; A91923; A91941; A91938; B29559
R:Kitamura, N.; Takagaki, Y.; Furuto, S.; Tanaka, T.; Nawa, H.; Nakanishi, S.
Nature 305, 545-549, 1983
A:Title: A single gene for bovine high molecular weight and low molecular weight kininogen
A:Reference number: A93317; PMID:84014106; PMID:6571699
A:Accession: A01282
A:Molecule type: mRNA
A:Residues: 1-619 <KT>
A:Cross-references: GB:V01492; GB:K01758; NID:9493; PIDN:CAA24736.1; PID:9494
R:Kato, H.; Nagasawa, S.; Suzuki, T.
J. Biochem. 67, 313-323, 1970
A:Title: Studies on the structure of bovine kininogen: cleavages of disulfide bonds and
A:Reference number: A91923; PMID:70180420; PMID:4986212
A:Accession: A91923
A:Molecule type: protein
A:Residues: 378-391 <KAT>
R:Han, Y.N.; Kato, H.; Iwanaga, S.; Suzuki, T.
J. Biochem. 79, 1201-1222, 1976
A:Title: Primary structure of bovine plasma high-molecular-weight kininogen. The amino a
A:Reference number: A91941; PMID:76260155; PMID:956151
A:Accession: A91941
A:Molecule type: protein
A:Residues: 387-455 <HAN>
A:Note: 398-Pro, 401-Val, and 455-Lys were also found
R:Han, Y.N.; Komiya, M.; Iwanaga, S.; Suzuki, T.
J. Biochem. 77, 55-68, 1975
A:Title: Studies on the primary structure of bovine high-molecular-weight kininogen. Ami
A:Reference number: A91938; PMID:75170265; PMID:1169237
A:Accession: A91938
A:Molecule type: protein
A:Residues: 456-496 <HA2>
R:Sueyoshi, T.; Miyata, T.; Hashimoto, N.; Kato, H.; Hayashida, H.; Miyata, T.; Iwanaga,
J. Biol. Chem. 262, 2768-2779, 1987
A:Title: Bovine high molecular weight kininogen. The amino acid sequence, positions of c
A:Reference number: A92627; PMID:87137530; PMID:3546295
A:Accession: B29559
A:Molecule type: protein
A:Residues: 'Z', 20-104, 'E', 106-256, 'XX', 257-376 <SUB>
R:Lottspeich, F.; Kellermann, J.; Henschen, A.; Foerzsch, B.; Muller-Esterl, W.
Eur. J. Biochem. 152, 307-314, 1985
A:Title: The amino acid sequence of the light chain of human high-molecular-mass kininog
A:Reference number: A91153; PMID:86030270; PMID:4054110
A:Contents: annotation; bovine cleavage sites; bovine carbohydrate binding sites
R:Sueyoshi, T.; Miyata, T.; Kato, H.; Iwanaga, S.
Seikagaku 56, 808, 1984
A:Title: Disulfide bonds in bovine HMW kininogens.
A:Reference number: A94300
A:Contents: annotation; disulfide bonds
A:Note: article in Japanese
C:Comment: the HMW kininogen precursor is produced from the same gene as the LMW form as
C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C:Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is impo
C:Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, i
xyproline residue is present in the kininogen prior to the release of bradykinin.
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupl
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-619/Product: HMW kininogen II #status predicted <MAT>
F:19-376/Product: HMW kininogen II heavy chain #status experimental <HCH>
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:261-372/Domain: cystatin homology <CY3>
F:377-386/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>

F:378-386/Product: bradykinin (kallidin I) #status experimental <BDY>
F:387-619/Product: HMW kininogen II light chain #status experimental <LCH>
F:418-488/Region: glycine/histidine/lysine-rich
F:19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status experimen
F:27-589, 82-93, 106-125, 141-144, 205-217, 228-247, 261-264, 325-337, 348-367/Disulfide bonds:
F:47/Binding site: carbohydrate (Asn) (covalent) #status absent
F:87, 168, 169, 204, 280/Binding site: carbohydrate (Asn) (covalent) #status experimental
F:136/Binding site: carbohydrate (Thr) (covalent) (partial) #status experimental
F:197/Binding site: carbohydrate (Asn) (covalent) (partial) #status experimental
F:376-377/Cleavage site: Met-Lys (kallikrein) #status experimental
F:380/Modified site: 4-hydroxyproline (Pro) #status predicted
F:386-387/Cleavage site: Arg-Ser (kallikrein) #status experimental
F:396, 400, 404, 510/Binding site: carbohydrate (Ser) (covalent) #status experimental
F:397, 398, 518, 522, 534, 546, 551, 569/Binding site: carbohydrate (Thr) (covalent) #status ex
F:496-497/Cleavage site: Arg-Thr (kallikrein) #status experimental

Query Match 68.2%; Score 450; DB 1; Length 619;

Best Local Similarity 68.9%; Pred. No. 5.1e-35;

Matches 84; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

QY 1 GKDFVOPPTKICVGCPRDPTNSPELEETLTHITIKLNAENNATFYFKIDNVKKARQVY 60

Db 252 GEDFL--PMWCVGCPKP:PVDSPLDEALNHSIAKLNAEHDTFYFKIDTVKKATVQVY 309

QY 61 AGKVFIDFVARETTCKESNEELTESCTKLGQSLDCNAEYVVPWEKKIYPTVNCOP 120

Db 310 GGLKYSIVFIARETTCKSGNEELTKSCININGQLLHCDANVYVVPWEKKYPTVNCOP 369

QY 121 LG 122

Db 370 LG 371

RESULT 7

A28055

K-kininogen, LMW I precursor - rat

C:Species: Rattus norvegicus (Norway rat)

C>Date: 20-Jun-1989 #sequence_revision 20-Jun-1989 #text_change 15-Nov-1996

C:Accession: A28055

R:Furuto-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.

J. Biol. Chem. 260, 12054-12059, 1985
A:Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and
inhibitor.

A:Reference number: A92496; PMID:86008264; PMID:2413018

A:Accession: A28055

A:Molecule type: mRNA

A:Residues: 1-433 <FUR>

C:Superfamily: kininogen; cystatin homology

C:Keywords: alternative splicing

F:1-18/Domain: signal sequence #status predicted <SIG>

F:19-433/Product: K-kininogen, LMW I #status predicted <MAT>

F:19-131/Domain: cystatin homology <CY1>

F:142-253/Domain: cystatin homology <CY2>

F:264-375/Domain: cystatin homology <CY3>

Query Match 64.5%; Score 426; DB 2; Length 433;

Best Local Similarity 65.0%; Pred. No. 6.6e-33;

Matches 80; Conservative 14; Mismatches 29; Indels 0; Gaps 0;

QY 1 GKDFVOPPTKICVGCPRDPTNSPELEETLTHITIKLNAENNATFYFKIDNVKKARQVY 60

Db 253 GDDLFEPLPDCPCPRNIPVDSPELKEALGHSIALNENHTFYFKIDTVKKATSOVY 312

QY 61 AGKVFIDFVARETTCKESNEELTESCTKLGQSLDCNAEYVVPWEKKIYPTVNCOP 120

Db 313 AGTKVIEFIARETTCKSKSNBELTADCTCKLQSLNCNANVYMPWENKVVPVTKCVY 372

QY 121 LGM 123

Db 373 LDM 375

RESULT 8

A25486
kininogen, HMW I precursor - rat
N:Contains: bradykinin
C:Species: Rattus norvegicus (Norway rat)
C>Date: 08-Mar-1989 #sequence_revision 08-Mar-1989 #text_change 15-Nov-1996
C:Accession: A25486
R:Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 262, 2190-2198, 1987
A:Title: Differing expression patterns and evolution of the rat kininogen gene family.
A:Reference number: A92625; MUID:87137443; PMID:3029068
A:Accession: A25486
A:Molecule type: mRNA
A:Residues: 1-639 <KIT>
A>Note: the authors translated the codon CAA for residue 347 as Asn
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-639/Product: kininogen, HMW I #status predicted <MAT>
F:19-131/Domain: cystatin homology <CY1>
F:142-253/Domain: cystatin homology <CY2>
F:264-375/Domain: cystatin homology <CY3>

Query Match 64.5%; Score 426; DB 2; Length 639;
Best Local Similarity 65.0%; Pred. No. 1e-32;
Matches 80; Conservative 14; Mismatches 29; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLAENNATFYKIDNVKKARVQV 60
Db 253 GDDLFSLLPKFCGCPKNIPVDSPELKEALGHSIAQLNAQNHLYFKIDTVKKATSQV 312

QY 61 AGKYYFIDFVARETTCSKESNEELTESCTKLGOSLDCNAEVYVVPWEKKIYPTVNCQP 120
Db 313 AGTKYVIEFIARETNCSTQNTLTADCTKHLGOSLNCNANVYRPNWKNVVPVTRCQA 372

QY 121 LGM 123
Db 373 LDM 374

RESULT 9
Major acute phase alpha-1 protein (version 2) - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 19-May-1989 #sequence_revision 19-May-1989 #text_change 20-Aug-1999
C:Accession: A23897; B23897
R:Anderson, K.P.; Heath, B.C.
J. Biol. Chem. 260, 12065-12071, 1985
A:Title: The relationship between rat major acute phase protein and the kininogens.
A:Reference number: A23897; MUID:86008266; PMID:2413019
A:Accession: A23897
A:Molecule type: protein
A:Residues: 1-14 <AND1>
A:Accession: B23897
A:Molecule type: mRNA
A:Residues: 5-430 <AND2>
A:Cross-references: GB:M11661; NID:G205307; PIDN:AAA41570.1; PID:G205308
A>Note: the authors translated the codon CTC for residue 410 as Arg, CTA for residue 415
C:Superfamily: kininogen; cystatin homology
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:263-374/Domain: cystatin homology <CY3>

Query Match 62.0%; Score 409; DB 2; Length 430;
Best Local Similarity 61.8%; Pred. No. 2.7e-31;
Matches 76; Conservative 15; Mismatches 32; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLAENNATFYKIDNVKKARVQV 60
Db 252 GDDLFSLLPKFCGCPKNIPVDSPELKEALGHSIAQLNAQNHLYFKIDTVKKATSQV 311

QY 61 AGKYYFIDFVARETTCSKESNEELTESCTKLGOSLDCNAEVYVVPWEKKIYPTVNCQP 120
Db 312 AGTKYVIEFIARETNCSTQNTLTADCTKHLGOSLNCNANVYRPNWKNVVPVTRCQA 371

QY 121 LGM 123
Db 372 LDM 374

RESULT 10
B28055
T-kininogen, LMW II precursor - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 20-Jun-1989 #sequence_revision 20-Jun-1989 #text_change 12-Dec-1997
C:Accession: B28055; E25486; E28526; C28526
R:Furuto-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.
J. Biol. Chem. 260, 12054-12059, 1985
A:Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and inhibitor.
A:Reference number: A92496; MUID:86008264; PMID:2413018
A:Accession: B28055
A:Molecule type: mRNA
A:Residues: 1-430 <FUR>
R:Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 262, 2190-2198, 1987
A:Title: Differing expression patterns and evolution of the rat kininogen gene family.
A:Reference number: A92625; MUID:87137443; PMID:3029068
A:Accession: E25486
A:Molecule type: DNA
A:Residues: 375-430 <KIT>
R:Enjoji, K.; Kato, H.; Hayaashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 973-979, 1988
A:Title: Purification and characterization of rat T-kininogens isolated from plasma of a
A:Reference number: A92729; MUID:88087226; PMID:3121623
A:Accession: B28526
A:Molecule type: protein
A:Residues: 'E', 20-25, 'MD', 28-48, 376-430 <ENJ>
A:Accession: C28526
A:Molecule type: protein
A:Residues: 'E', 20-48, 376-388, 'R', 390-419, 'ER', 422-430 <EN2>
C:Superfamily: kininogen; cystatin homology
C:Keywords: glycoprotein; pyroglutamic acid
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-430/Product: T-kininogen, LMW II #status experimental <MAT>
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:263-374/Domain: cystatin homology <CY3>
F:19/Modified site: pyroglutamic acid (Gln) (in mature form) #status experiment
F:82,126,168,204,326/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:83-94,107-125,141-144,205-217,228-247,263-286,327-339,350-369/Disulfide bonds: #status

Query Match 62.0%; Score 409; DB 2; Length 430;
Best Local Similarity 61.8%; Pred. No. 2.7e-31;
Matches 76; Conservative 15; Mismatches 32; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITIKLAENNATFYKIDNVKKARVQV 60
Db 252 GDDLFSLLPKFCGCPKNIPVDSPELKEALGHSIAQLNAQNHLYFKIDTVKKATSQV 311

QY 61 AGKYYFIDFVARETTCSKESNEELTESCTKLGOSLDCNAEVYVVPWEKKIYPTVNCQP 120
Db 312 AGTKYVIEFIARETNCSTQNTLTADCTKHLGOSLNCNANVYRPNWKNVVPVTRCQA 371

QY 121 LGM 123
Db 372 LDM 374

RESULT 11
KGRFM
Major acute phase alpha-1 protein precursor - rat (fragment)
N:Contains: bradykinin
C:Species: Rattus norvegicus (Norway rat)
C>Date: 27-Nov-1985 #sequence_revision 27-Nov-1985 #text_change 12-Apr-1996
C:Accession: A01285
R:Coile, T.; Inglis, A.S.; Roxburgh, C.M.; Howlett, G.J.; Schreiber, G.

FEBS Lett. 182, 57-61, 1985
A:Title: Major acute phase alpha-1-protein of the rat is homologous to bovine kininogen A
A:Reference number: A01285; MUID:85127561; PMID:2578992
A:Accession: A01285
A:Molecule type: mRNA
A:Residues: 1-423 <COL>
C:Comment: This plasma glycoprotein inhibits cysteine proteinases. During acute inflammation
X:
C:Superfamily: kininogen; cystatin homology
C:Keywords: bradykinin; cysteine proteinase inhibitor; duplication; glycoprotein; inflammation
F:1-11/Domain: signal sequence (fragment) #status predicted <SIG>
F:12-423/Product: major acute phase alpha-1 protein #status predicted <MAP>
F:12-123/Domain: cystatin homology <CY1>
F:134-245/Domain: cystatin homology <CY2>
F:256-367/Domain: cystatin homology <CY3>
F:371-379/Product: bradykinin #status predicted <BDY>
F:12/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status predicted
F:161,197/Binding site: carbohydrate (Asn) (covalent) #status predicted
Query Match 60.9%; Score 402; DB 1; Length 423;
Best Local Similarity 61.8%; Pred. No. 1.2e-30;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;
Qy 1 GKDFVQPTKICVGCPRDIPNSPELEETLTHITTKNAENNAFFYFKIDNVKARVQV 60
Db 245 GDDLPELLPKNCRGCPREIPVDSPELKEALGHSIARLNQNHIFYPKIDTVKATSQV 304
Qy 61 AGKYFIDFVARETTCKSESNEELTESCTKKLGSLDCNAEVVVPWEKKIVPTVNCOP 120
Db 305 AGVIYVIEFIARETNCQSKTELTADCTCKHLGSLNCNANVYRPNENKVPTVRCQA 364
Qy 121 LGM 123
Db 365 LDM 367
RESULT 12
KGRTTI
N:Alternate names: 73K protein; LMW kininogen T-I
N:Contains: bradykinin; T-kinin
C:Species: Rattus norvegicus (Norway rat)
C:Date: 17-Mar-1987 #sequence revision 17-Mar-1987 #text change 22-Jun-1999
C:Accession: A01286; D25486; E00193; JQ0027; E25488; A28525; S68036
R:Furuto-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.
J. Biol. Chem. 260, 12054-12059, 1985
A:Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and inhibitor.
A:Reference number: A92496; MUID:86008264; PMID:2413018
A:Accession: A01286
A:Molecule type: mRNA
A:Residues: 1-430 <FUR>
A:Cross-references: GB:M11883; NID:g205084; PIDN:AAA41489.1; PID:g205085
R:Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 262, 2190-2198, 1987
A:Title: Differing expression patterns and evolution of the rat kininogen gene family.
A:Reference number: A32625; MUID:87137443; PMID:3029068
A:Accession: D25486
A:Molecule type: DNA
A:Residues: 375-430 <KIT>
R:Eniyoji, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 973-979, 1988
A:Title: Purification and characterization of rat T-kininogens isolated from plasma of a
A:Reference number: A92729; MUID:88087226; PMID:3121623
A:Accession: A28526
A:Molecule type: protein
A:Residues: 'E', 20-48; 376-430 <BNJ>
R:Kanda, S.; Sugiyama, K.; Takahashi, M.; Shumiya, S.; Tomino, S.; Nagase, S.
Jpn. J. Cancer Res. 81, 63-68, 1990
A:Title: Identification of a protein increasing in serum of Nagase analbuminemic rats be
A:Reference number: PLO193; MUID:90216390; PMID:2108948
A:Accession: PLO193
A:Molecule type: mRNA

A:Residues: 330-420, 'R', 422-429, 'P' <KAN>
R:Anderson, K.P.; Croyle, M.L.; Lingrel, J.B.
Gene 81, 119-128, 1989
A:Title: Primary structure of a gene encoding rat T-kininogen.
A:Reference number: JQ0027; MUID:90034172; PMID:2806908
A:Accession: JQ0027
A:Molecule type: DNA
A:Residues: 1-60, 'E', 62-113, 'R', 115-165, 'F', 167-178, 'TKI', 182-211, 'F', 213-256, 'S', 259-388
A:Experimental source: strain Sprague-Dawley
R:Kagayama, R.; Kitamura, N.; Okubo, H.; Nakanishi, S.
J. Biol. Chem. 262, 2345-2355, 1987
A:Title: Differing utilization of homologous transcription initiation sites of rat K and
A:Reference number: A25488; MUID:87137465; PMID:3818598
A:Accession: B25488
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-48 <XAG>
A:Cross-references: GB:M14356; NID:g205090; PIDN:AAA41492.1; PID:g205091
R:Eniyoji, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
J. Biol. Chem. 263, 965-972, 1988
A:Title: Purification and characterization of two kinds of low molecular weight kininogen
A:Reference number: A28525; MUID:88087225; PMID:3335530
A:Accession: A28525
A:Molecule type: protein
A:Residues: 376-430 <EN2>
R:Sierra, F.; Walter, R.; Vautravers, P.; Guigoz, Y.
Arch. Biochem. Biophys. 322, 333-338, 1995
A:Title: Identification of several isoforms of T-kininogen expressed in the liver of agi
A:Reference number: S68034; MUID:96032652; PMID:7574705
A:Accession: S68036
A:Molecule type: mRNA
A:Residues: 340-430 <SIE>
A:Experimental source: clone pSG17
C:Comment: At least three types of LMW kininogen precursors are present in rat plasma, t
ceding bradykinin.
C:Comment: T-kininogens contain T-kinin (I-S-bradykinin), a novel kinin isolated after t
d of an Arg or Lys, it is probably not released from its precursor by either tissue or p
C:Comment: The T-kininogens are produced in response to an inflammatory stimulant.
C:Genetics:
A:Introns: 65/3; 102/3; 130/1; 187/3; 223/2; 252/1; 309/3; 345/3; 374/3; 398/3
C:Superfamily: kininogen; cystatin homology
C:Keywords: acute phase; bradykinin; cysteine proteinase inhibitor; duplication; glycop
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-430/Product: T-kininogen I #status experimental <MAT>
F:19-130/Domain: cystatin homology <CY1>
F:141-252/Domain: cystatin homology <CY2>
F:263-374/Domain: cystatin homology <CY3>
F:378-386/Product: bradykinin #status predicted <BDY>
F:19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status experimen
F:82, 126, 168, 204, 326/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:83-94, 107-125, 141-144, 205-217, 228-247, 263-266, 327-339, 350-369/Disulfide bonds: #status

Query Match 60.8%; Score 401; DB 1; Length 430;
Best Local Similarity 61.8%; Pred. No. 1.6e-30;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

Qy 1 GKDFVQPTKICVGCPRDIPNSPELEETLTHITTKNAENNAFFYFKIDNVKARVQV 60

Db 252 GDDLPELLPKNCRGCPREIPVDSPELKEALGHSIARLNQNHIFYPKIDTVKATSQV 311

Qy 61 AGKYFIDFVARETTCKSESNEELTESCTKKLGSLDCNAEVVVPWEKKIVPTVNCOP 120

Db 312 AGVIYVIEFIARETNCQSKTELTADCTCKHLGSLNCNANVYRPNENKVPTVRCQA 371

Qy 121 LGM 123

Db 372 LDM 374

RESULT 13

UDBO

Cyscatin - bovine
N:Alternate names: thiol proteinase inhibitor

C:Species: Bos primigenius taurus (cattle)
C:Date: 28-Feb-1986 #sequence_revision 28-Feb-1986 #text_change 06-Dec-1996
C:Accession: A01271
R: Hirado, M.; Tsunawawa, S.; Sakiyama, F.; Niinobe, M.; Fujii, S.
FEBS Lett. 186, 41-45, 1985
A:Title: Complete amino acid sequence of bovine colostrum low-M-r cysteine proteinase in
A:Reference number: A01271; MUID:85231205; PMID:3891407
A:Accession: A01271

A:Molecule type: protein
A:Residues: 1-112 <HIR>
C:Superfamily: cystatin; cystatin homology
C:Keywords: colostrum; cysteine proteinase inhibitor
F:2-112/Domain: cystatin homology <CYS>
F:48-52/Region: inhibitory #status predicted
F:86-76,90-110/Disulfide bonds: #status predicted

Query Match 20.6%; Score 136; DB 1; Length 112;
Best Local Similarity 30.6%; Pred. No. 6.4e-06;
Matches 34; Conservative 22; Mismatches 31; Indels 24; Gaps 5;

QY 22 NSPELEETITHTIKLNAENNAFFPKIDNVKARVQVAGKYYFIDFVARETTCSKSN 81

DB 12 NEEGQVEALSFVAFSEFNKSNDAIOSRVVVRVVRARQVQVGMNYPDLVELGRITCTK--S 69

QY 82 BELTESC-----ETKKLGQSLDCNAEVVVPWEKKIYPTVN-----CQ 119

DB 70 QANLDSCPHNPQHLKREK-----CSFQVYVVPWMN-----TINLVKFSQ 111

RESULT 14

S68034

T-kininogen (clone pSG22) - rat (fragment)
C:Species: Rattus norvegicus (Norway rat)
C:Date: 12-Feb-1998 #sequence_revision 20-Feb-1998 #text_change 07-May-1999
C:Accession: S68034
R: Sierra, F.; Walter, R.; Vautravers, P.; Guigoz, Y.
Arch. Biochem. Biophys. 322, 333-338, 1995
A:Title: Identification of several isoforms of T-kininogen expressed in the liver of ag
A:Reference number: S68034; MUID:96032652; PMID:7574705
A:Accession: S68034

A:Molecule type: mRNA

A:Residues: 1-91 <SIE>

A:Experimental source: liver

C:Superfamily: kininogen; cystatin homology

C:Keywords: alternative splicing; cysteine proteinase inhibitor; plasma

Query Match 20.2%; Score 133; DB 2; Length 91;
Best Local Similarity 68.6%; Pred. No. 9.8e-06;
Matches 24; Conservative 3; Mismatches 9; Indels 0; Gaps 0;

QY 89 ETKKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 123

DB 1 ETKKLGQSLDCNAEVVVPWEKKIYPTVNCQPLDM 35

RESULT 15

S68035

T-kininogen (clone pSG17) - rat (fragment)
C:Species: Rattus norvegicus (Norway rat)
C:Date: 12-Feb-1998 #sequence_revision 20-Feb-1998 #text_change 07-May-1999
C:Accession: S68035
R: Sierra, F.; Walter, R.; Vautravers, P.; Guigoz, Y.
Arch. Biochem. Biophys. 322, 333-338, 1995
A:Title: Identification of several isoforms of T-kininogen expressed in the liver of ag
A:Reference number: S68034; MUID:96032652; PMID:7574705
A:Accession: S68035

A:Molecule type: mRNA

A:Residues: 1-91 <SIE>

A:Experimental source: liver

C:Superfamily: kininogen; cystatin homology

C:Keywords: alternative splicing; cysteine proteinase inhibitor; plasma

Query Match 20.2%; Score 133; DB 2; Length 91;

Best Local Similarity 68.6%; Pred. No. 9.8e-06;
Matches 24; Conservative 3; Mismatches 8; Indels 0; Gaps 0;

QY 89 ETKKLGQSLDCNAEVVVPWEKKIYPTVNCQPLGM 123

DB 1 ETKKLGQSLDCNAEVVVPWEKKIYPTVNCQPLDM 35

Search completed: September 24, 2004, 14:10:48
Job time : 14.284 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:08:41 : Search time 43.296 Seconds
(without alignments)
913.519 Million cell updates/sec

Title: US-10-661-784-1
Perfect score: 660
Sequence: 1 GKDFVQPTKICVGRDIP.....YVWPWEKKIYPTNCQPLGM 123

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1349238 seqs, 321558718 residues

Total number of hits satisfying chosen parameters: 1349238

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications AA:*
1: /cgn2_6/ptodata/1/pubpaa/US07_PUBCOMB.pep.*
2: /cgn2_6/ptodata/1/pubpaa/PCT_NEW_PUB.pep.*
3: /cgn2_6/ptodata/1/pubpaa/US06_NEW_PUB.pep.*
4: /cgn2_6/ptodata/1/pubpaa/US06_PUBCOMB.pep.*
5: /cgn2_6/ptodata/1/pubpaa/US07_NEW_PUB.pep.*
6: /cgn2_6/ptodata/1/pubpaa/PCTUS_PUBCOMB.pep.*
7: /cgn2_6/ptodata/1/pubpaa/US08_NEW_PUB.pep.*
8: /cgn2_6/ptodata/1/pubpaa/US08_PUBCOMB.pep.*
9: /cgn2_6/ptodata/1/pubpaa/US09A_PUBCOMB.pep.*
10: /cgn2_6/ptodata/1/pubpaa/US09B_PUBCOMB.pep.*
11: /cgn2_6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubpaa/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	560	100.0	390	15	US-10-162-335-82
2	560	100.0	398	15	US-10-162-335-70
3	560	100.0	427	10	US-09-919-039-29
4	560	100.0	615	15	US-10-162-335-72
5	560	100.0	644	15	US-10-162-335-74
6	560	100.0	624	15	US-10-162-335-84
7	402	60.9	424	14	US-10-316-253-217
8	401	60.8	430	14	US-10-316-253-215
9	169	25.6	178	9	US-09-969-834-1
10	165.5	25.1	145	14	US-10-329-428-2
11	165.5	25.1	167	10	US-09-746-783-197
12	138.5	21.0	121	9	US-09-775-932-14
13	138.5	21.0	128	9	US-09-775-932-12
14	138.5	21.0	149	9	US-09-940-497-2
15	136	20.6	112	8	US-08-849-303-16

Sequence 16, Appl
Sequence 24, Appl
Sequence 19, Appl
Sequence 19, Appl
Sequence 46, Appl
Sequence 48, Appl
Sequence 82, Appl
Sequence 86, Appl
Sequence 84, Appl
Sequence 78, Appl
Sequence 80, Appl
Sequence 93, Appl
Sequence 425, Appl
Sequence 26, Appl
Sequence 26, Appl
Sequence 2, Appl
Sequence 18, Appl
Sequence 18, Appl
Sequence 17, Appl
Sequence 3, Appl
Sequence 3, Appl
Sequence 3, Appl
Sequence 47, Appl
Sequence 17, Appl
Sequence 1, Appl
Sequence 4, Appl
Sequence 2, Appl
Sequence 8, Appl
Sequence 24, Appl

ALIGNMENTS

RESULT 1
US-10-162-335-82
; Sequence 82, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalte, Tor
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Miller, Isabelle
; APPLICANT: Padigaru, Muraidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shimkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Sytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Zoss, Edward Z.
; APPLICANT: Aarhusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 82
; LENGTH: 390
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-82

Query Match 100.0%; Score 660; DB 15; Length 390;
Best Local Similarity 100.0%; Pred. No. 2.9e-64;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GKDVPPTKICVGCPRDPTSPLEELTHITTKLAENNATYFKIDNVKARQVW 60
Db 216 GKDVPPTKICVGCPRDPTSPLEELTHITTKLAENNATYFKIDNVKARQVW 275

Qy 61 AGKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEVYVPWEKKIYPTVNCOP 120
Db 276 AGKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEVYVPWEKKIYPTVNCOP 335

Qy 121 LGM 123
Db 336 LGM 338

RESULT 2
US-10-162-335-70
; Sequence 70, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Bainger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjal, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malyankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigaru, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shinkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; APPLICANT: Zerhusen, Bryan D.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B

; CURRENT APPLICATION NUMBER: US/10/162,335
; CURRENT FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 70
; LENGTH: 398
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-70

Query Match 100.0%; Score 660; DB 15; Length 398;
Best Local Similarity 100.0%; Pred. No. 3e-64; 0; Indels 0; Gaps 0;
Matches 123; Conservative 0; Mismatches 0;

Qy 1 GKDVPPTKICVGCPRDPTSPLEELTHITTKLAENNATYFKIDNVKARQVW 60
Db 224 GKDVPPTKICVGCPRDPTSPLEELTHITTKLAENNATYFKIDNVKARQVW 283

Qy 61 AGKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEVYVPWEKKIYPTVNCOP 120
Db 284 AGKYFIDFVARETTCKESNEELTESCETKLGQSLDCNAEVYVPWEKKIYPTVNCOP 343

Qy 121 LGM 123
Db 344 LGM 346

RESULT 3
US-09-919-039-29
; Sequence 29, Application US/09919039
; Publication No. US20030108871A1
; GENERAL INFORMATION:
; APPLICANT: Kaser, Matthew R.
; TITLE OF INVENTION: GENES EXPRESSED IN TREATED HUMAN C3A LIVER CELL CULTURES
; FILE REFERENCE: PA-0035 US
; CURRENT APPLICATION NUMBER: US/09/919,039
; CURRENT FILING DATE: 2002-09-09
; PRIOR APPLICATION NUMBER: 60/222,113
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 401
; SOFTWARE: PERL Program
; SEQ ID NO 29
; LENGTH: 427
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. US20030108871A1 167507CD1
US-09-919-039-29

Query Match 100.0%; Score 660; DB 10; Length 427;
Best Local Similarity 100.0%; Pred. No. 3.3e-64;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;


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QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKARQVW 60
Db 253 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKARQVW 312
QY 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTVNCOP 120
Db 313 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTVNCOP 372
QY 121 LGM 123
Db 373 LGM 375

RESULT 4
US-10-162-335-72
; Sequence 72, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalt, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malvankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigar, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shinkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; PRIOR FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 72
; LENGTH: 615
; TYPE: PRT
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; ORGANISM: Homo sapiens
US-10-162-335-72
Query Match 100.0%; Score 660; DB 15; Length 615;
Best Local Similarity 100.0%; Pred. No. 5.4e-64; Indels 0; Gaps 0;
Matches 123; Conservative 0; Mismatches 0;
QY 1 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKARQVW 60
Db 224 GKDFVQPTKICVGCPRDIPNTSPLEETLTHITKLNAENNATFYFKIDNVKARQVW 283
QY 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTVNCOP 120
Db 284 AGKKYFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTVNCOP 343
QY 121 LGM 123
Db 344 LGM 346

RESULT 5
US-10-162-335-72
; Sequence 74, Application US/10162335
; Publication No. US20040009480A1
; GENERAL INFORMATION:
; APPLICANT: Anderson, David W.
; APPLICANT: Baumgartner, Jason C.
; APPLICANT: Boldog, Ferenc L.
; APPLICANT: Casman, Stacie J.
; APPLICANT: Edinger, Shlomit R.
; APPLICANT: Gangolli, Esha A.
; APPLICANT: Gerlach, Valerie
; APPLICANT: Gorman, Linda
; APPLICANT: Guo, Xiaojia (Sasha)
; APPLICANT: Hjalt, Tord
; APPLICANT: Kekuda, Ramesh
; APPLICANT: Li, Li
; APPLICANT: MacDougall, John R.
; APPLICANT: Malvankar, Uriel M.
; APPLICANT: Millet, Isabelle
; APPLICANT: Padigar, Muralidhara
; APPLICANT: Patturajan, Meera
; APPLICANT: Pena, Carol E. A.
; APPLICANT: Rastelli, Luca
; APPLICANT: Shinkets, Richard A.
; APPLICANT: Stone, David J.
; APPLICANT: Spytek, Kimberly A.
; APPLICANT: Vernet, Corine A. M.
; APPLICANT: Voss, Edward Z.
; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
; FILE REFERENCE: 21402-377 B
; CURRENT APPLICATION NUMBER: US/10/162,335
; PRIOR FILING DATE: 2002-10-01
; PRIOR APPLICATION NUMBER: 60/295,607
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/295,661
; PRIOR FILING DATE: 2001-06-04
; PRIOR APPLICATION NUMBER: 60/296,404
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/296,418
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: 60/297,414
; PRIOR FILING DATE: 2001-06-11
; PRIOR APPLICATION NUMBER: 60/297,567
; PRIOR FILING DATE: 2001-06-12
; PRIOR APPLICATION NUMBER: 60/298,285
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
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; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 74
; LENGTH: 644
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-74

Query Match 100.0%; Score 660; DB 15; Length 644;
Best Local Similarity 100.0%; Pred. No. 5.7e-64;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 60
Db 253 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 312
Qy 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVYVVPWEKKIYPTVNCOP 120
Db 313 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVYVVPWEKKIYPTVNCOP 372
Qy 121 LGM 123
Db 373 LGM 375

RESULT 6

US-10-162-335-84

; Sequence 84, Application US/10162335

; Publication No. US20040009480A1

; GENERAL INFORMATION:

; APPLICANT: Anderson, David W.

; APPLICANT: Baumgartner, Jason C.

; APPLICANT: Boldog, Ferenc L.

; APPLICANT: Casman, Stacie J.

; APPLICANT: Edinger, Shlomit R.

; APPLICANT: Gangolli, Esha A.

; APPLICANT: Gerlach, Valerie

; APPLICANT: Gorman, Linda

; APPLICANT: Guo, Xiaojia (Sasha)

; APPLICANT: Hjalte, Tord

; APPLICANT: Kekuda, Ramesh

; APPLICANT: Li, Li

; APPLICANT: MacDougall, John R.

; APPLICANT: Malvankar, Uriel M.

; APPLICANT: Millet, Isabelle

; APPLICANT: Padigaru, Muralidhara

; APPLICANT: Patturajan, Meera

; APPLICANT: Pena, Carol E. A.

; APPLICANT: Rastelli, Luca

; APPLICANT: Shimkets, Richard A.

; APPLICANT: Stone, David J.

; APPLICANT: Spytek, Kimberly A.

; APPLICANT: Vernet, Corine A. M.

; APPLICANT: Voss, Edward Z.

; APPLICANT: Zerhusen, Bryan D.

; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method

; FILE REFERENCE: 21402-377 B

; CURRENT APPLICATION NUMBER: US/10/162,335

; CURRENT FILING DATE: 2002-10-01

; PRIOR APPLICATION NUMBER: 60/295,607

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/295,661

; PRIOR FILING DATE: 2001-06-04

; PRIOR APPLICATION NUMBER: 60/296,404

; PRIOR FILING DATE: 2001-06-06

; PRIOR APPLICATION NUMBER: 60/296,418

; PRIOR FILING DATE: 2001-06-06

; PRIOR APPLICATION NUMBER: 60/297,414

; PRIOR FILING DATE: 2001-06-11

; PRIOR APPLICATION NUMBER: 60/297,567

; PRIOR FILING DATE: 2001-06-12

; PRIOR APPLICATION NUMBER: 60/298,285

; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: 60/298,556
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: 60/299,949
; PRIOR FILING DATE: 2001-06-21
; PRIOR APPLICATION NUMBER: 60/300,883
; PRIOR FILING DATE: 2001-06-26
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 201
; SEQ ID NO 84
; LENGTH: 644
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-335-84

Query Match 100.0%; Score 660; DB 15; Length 644;
Best Local Similarity 100.0%; Pred. No. 5.7e-64;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 60
Db 253 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 312
Qy 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVYVVPWEKKIYPTVNCOP 120
Db 313 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVYVVPWEKKIYPTVNCOP 372
Qy 121 LGM 123
Db 373 LGM 375

RESULT 7

US-10-316-253-217

; Sequence 217, Application US/10316253

; Publication No. US20030162706A1

; GENERAL INFORMATION:

; APPLICANT: The Procter & Gamble Company

; APPLICANT: Peters, Kevin

; APPLICANT: Thompson, Larry

; APPLICANT: Wang, Feng

; APPLICANT: Greis, Kenneth

; TITLE OF INVENTION: Angiogenesis Modulating Proteins

; FILE REFERENCE: 8865M

; CURRENT APPLICATION NUMBER: US/10/316,253

; CURRENT FILING DATE: 2002-12-10

; PRIOR APPLICATION NUMBER: US 60/355,295

; PRIOR FILING DATE: 2002-02-08

; NUMBER OF SEQ ID NOS: 308

; SOFTWARE: Patent in version 3.1

; SEQ ID NO 217

; LENGTH: 424

; TYPE: PRT

; ORGANISM: Rattus norvegicus

US-10-316-253-217

Query Match 60.9%; Score 402; DB 14; Length 424;
Best Local Similarity 61.8%; Pred. No. 1.1e-35;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

Qy 1 GKDFVQPTKICVGCPRDPTNSPELEETLTHITKLAENNAATFYFKIDNVKKARQVQV 60
Db 246 GDDLPELLPKNKGCPREIPVDSPELKEALGHSLARLNQHNIHFYFKIDTVKATSOV 305
Qy 61 AGKKYFIDFVARETTCSKESNEELTESCETKLGQSLDCNAEVYVVPWEKKIYPTVNCOP 120
Db 306 AGVYVIEFIARETNCQSKTSLTADCTKHLGQSLNCNANVYMPFENKVVPTVRCOA 365
Qy 121 LGM 123
Db 366 LDM 368

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RESULT 8
US-10-316-253-215
; Sequence 215, Application US/10316253
; Publication No. US20030162706A1
; GENERAL INFORMATION:
; APPLICANT: The Procter & Gamble Company
; APPLICANT: Peters, Kevin
; APPLICANT: Thompson, Larry
; APPLICANT: Wang, Feng
; APPLICANT: Greis, Kenneth
; TITLE OF INVENTION: Angiogenesis Modulating Proteins
; FILE REFERENCE: 8665M
; CURRENT APPLICATION NUMBER: US/10/316,253
; CURRENT FILING DATE: 2002-12-10
; PRIOR APPLICATION NUMBER: US 60/355,295
; PRIOR FILING DATE: 2002-02-08
; NUMBER OF SEQ ID NOS: 308
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 215
; LENGTH: 430
; TYPE: PRT
; ORGANISM: Rattus norvegicus
US-10-316-253-215

Query Match 60.8%; Score 401; DB 14; Length 430;
Best Local Similarity 61.8%; Pred. No. 1.5e-35;
Matches 76; Conservative 14; Mismatches 33; Indels 0; Gaps 0;

QY 1 GKDFVQPTKICVGPDRDIPNPSPELEETLTHITKLAENNAATFYFKIDNVKARVQV 60
Db 252 GDDLPELLPKRCGCPREIPVDSPELKEALGHSIAQLNAQHNHIFYFKIDTVKKAQV 311

QY 61 AGKYFIDFVAREITCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVNCOP 120
Db 312 AGVIIVIEFIAREITCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVNCOP 371

QY 121 LGM 123
Db 372 LDM 374

RESULT 9
US-09-969-834-1
; Sequence 1, Application US/09969834
; Patent No. US20020102711A1
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESS: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/969,834
; FILING DATE: 01-Oct-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/09/471,765
; FILING DATE: <Unknown>
; APPLICATION NUMBER: US/08/791,522
; FILING DATE: <Unknown>
; APPLICATION NUMBER: 09/471,765

FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0193 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 178 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
CLONE: 30443
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-969-834-1

Query Match 25.6%; Score 169; DB 9; Length 178;
Best Local Similarity 32.5%; Pred. No. 2.1e-10;
Matches 39; Conservative 22; Mismatches 49; Indels 10; Gaps 4;

QY 9 TKICVGPDRDIPNPSPELEETLTHITKLAENNAATFYFKIDNVKARVQVWAGKYFID 69
Db 54 SRVKPGFPKTIKNDPGVLQAARYSVEKENNCTNDMFLFKESRITRALVQIVKGLKXMLE 113

QY 69 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVNCOP 123
Db 114 VEIGRTTCKKNQHLRL-DDCDFQTNHTLKQTLSCYSEVWVVPW---VPALRGACSPSL 168

RESULT 10
US-10-329-428-2
; Sequence 2, Application US/10329428
; Publication No. US20030114646A1
; GENERAL INFORMATION:
; APPLICANT: Li, et al.
; TITLE OF INVENTION: Human Cystatin F
; FILE REFERENCE: PF265PID2
; CURRENT APPLICATION NUMBER: US/10/329,428
; CURRENT FILING DATE: 2002-12-27
; PRIOR APPLICATION NUMBER: 60/014,795
; PRIOR FILING DATE: 1996-04-03
; PRIOR APPLICATION NUMBER: 08/832,535
; PRIOR FILING DATE: 1997-04-03
; PRIOR APPLICATION NUMBER: 09/019,485
; PRIOR FILING DATE: 1998-01-29
; PRIOR APPLICATION NUMBER: 09/528,436
; PRIOR FILING DATE: 2000-03-17
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 2
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-329-428-2

Query Match 25.1%; Score 165.5; DB 14; Length 145;
Best Local Similarity 32.5%; Pred. No. 3.9e-10;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;

QY 9 TKICVGPDRDIPNPSPELEETLTHITKLAENNAATFYFKIDNVKARVQVWAGKYFID 69
Db 32 SRVKPGFPKTIKNDPGVLQAARYSVEKENNCTNDMFLFKESRITRALVQIVKGLKXMLE 91

QY 69 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKI-YPTVNC 118
Db 92 VEIGRTTCKKNQHLRL-DDCDFQTNHTLKQTLSCYSEVWVVPWLPQHFVPLVLC 144

RESULT 11
US-09-746-783-197
```

; Sequence 197, Application US/09746783
; Publication No. US20030044935A1
; GENERAL INFORMATION:
; APPLICANT: Jacobs, Kenneth
; McCoy, John M.
; LaValle, Edward R.
; Racie, Lisa A.
; Treacy, Maurice
; Spaulding, Vikki
; Agostino, Michael J.
; Howes, Steven H.
; Fectel, Kim
; TITLE OF INVENTION: SECRETED PROTEINS AND POLYNUCLEOTIDES
; ENCODING THEM
; NUMBER OF SEQUENCES: 231
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genetics Institute, Inc.
; STREET: 87 Cambridgepark Drive
; CITY: Cambridge
; STATE: MA
; COUNTRY: U.S.A.
; ZIP: 02140
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/746,783
; FILING DATE: 21-Dec-2000
; CLASSIFICATION: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Milasincic, Debra J.
; REGISTRATION NUMBER: 46,931
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 227-7400
; TELEFAX: (617) 742-4214
; INFORMATION FOR SEQ ID NO: 197:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 167 amino acids
; TYPE: amino acid
; STRANDEDNESS: <Unknown>
; TOPOLOGY: linear
; MOLECULE TYPE: Protein
; SEQUENCE DESCRIPTION: SEQ ID NO: 197:
US-09-746-783-197

Query Match 25.1%; Score 165.5; DB 10; Length 167;
Best Local Similarity 32.5%; Pred. No. 4.7e-10;
Matches 37; Conservative 22; Mismatches 50; Indels 5; Gaps 3;

QY 9 TKICVCPDRDIPNPELEETLTHITKLNAENNAATFYFKIDNVKARVQVVGKYYFI 68
Db 54 SRVKGFPKTKINDPVGIAARYSVKFNCTNDMLFKESRIITRALVQIVKGLKYLE 113

QY 69 FVARETTCSKE---TKLGSQSLDCAEVVVPWEKKI-YPTVNC 118
Db 114 VEIGRTTCKNQHLRL-DDCDQTNHTLAQTILSCYSEVVVVPWLQHFVEFVLRC 166

RESULT 12
US-09-775-932-14
; Sequence 14, Application US/09775932
; Patent No. US20020137671A1
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503

; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 14
; LENGTH: 121
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-14

Query Match 21.0%; Score 138.5; DB 9; Length 121;
Best Local Similarity 31.5%; Pred. No. 3e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 8 PTKICVCPDRDIPNPELEETLTHITKLNAENNAATFYFKIDNVKARVQVVGKYYFI 67
Db 2 PQRWVGELRDLSPDDPQVQKAAQAAVASYNMGNSIYYFRDTHIIKAQSQLVAGIKYFL 61

QY 68 DFVARETTCSKE---SNEELTESCETKKLGO--SLDCNAEVVVPWE 109
Db 62 TMEVGSTDCRKTAVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 108

RESULT 13
US-09-775-932-12
; Sequence 12, Application US/09775932
; Patent No. US20020137671A1
; GENERAL INFORMATION:
; APPLICANT: University of British Columbia
; TITLE OF INVENTION: Production and use of Modified Cystatins
; FILE REFERENCE: 58069
; CURRENT APPLICATION NUMBER: US/09/775,932
; CURRENT FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: CA99/00717
; PRIOR FILING DATE: 1999-08-05
; PRIOR APPLICATION NUMBER: 60/095,503
; PRIOR FILING DATE: 1998-08-05
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 12
; LENGTH: 128
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-775-932-12

Query Match 21.0%; Score 138.5; DB 9; Length 128;
Best Local Similarity 31.5%; Pred. No. 3.2e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 8 PTKICVCPDRDIPNPELEETLTHITKLNAENNAATFYFKIDNVKARVQVVGKYYFI 67
Db 9 PQRWVGELRDLSPDDPQVQKAAQAAVASYNMGNSIYYFRDTHIIKAQSQLVAGIKYFL 68

QY 68 DFVARETTCSKE---SNEELTESCETKKLGO--SLDCNAEVVVPWE 109
Db 69 TMEVGSTDCRKTAVTGDHVDLT-TCPLAAGAQQEKLRCDFEVLVVPWQ 115

RESULT 14
US-09-940-497-2
; Sequence 2, Application US/09940497
; Patent No. US20020052476A1
; GENERAL INFORMATION:
; APPLICANT: Ni et al.
; TITLE OF INVENTION: Human Cystatin E
; FILE REFERENCE: PE202P1D2
; CURRENT APPLICATION NUMBER: US/09/940,497
; CURRENT FILING DATE: 2001-08-29
; PRIOR APPLICATION NUMBER: US 09/241,376
; PRIOR FILING DATE: 1999-02-02
; PRIOR APPLICATION NUMBER: US 08/744,138
; PRIOR FILING DATE: 1996-11-05
; PRIOR APPLICATION NUMBER: US 08/461,030
; PRIOR FILING DATE: 1995-06-05

; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 2
; LENGTH: 149
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-940-497-2

Query Match 21.0%; Score 138.5; DB 9; Length 149;
Best Local Similarity 31.5%; Pred. No. 3.9e-07;
Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

QY 8 PTKICVGPDRIPNTSPLEETLTHTITKLAENNATFYFKIDNVKKARVQVWAGKYFI 67

Db 30 PQRVMVGLRLDSPDPQVQKAAQAAVASYNMGNSIYFRDTHIIKQSLVAGIKYFL 89

QY 68 DFVARETTCSEK-----SNEELTESCETKKLGQ--SLDCNAEYVVPWE 109

Db 90 TMEGSTDCKRTRVTGDHVDLT-TCPLAAGAQCEKRLCDFEVLVVPWQ 136

RESULT 15

US-08-849-303-16
; Sequence 16, Application US/08849303
; Publication No. US20030221209A1

GENERAL INFORMATION:

; APPLICANT: Atkinson, Howard J.
; APPLICANT: McPherson, Michael J.
; APPLICANT: Urwin, Peter E.

; TITLE OF INVENTION: MODIFIED PROTEINASE INHIBITORS

; NUMBER OF SEQUENCES: 79

; CORRESPONDENCE ADDRESS:

; ADDRESS: Klauber & Jackson

; STREET: 411 Hackensack Avenue, 4th Floor

; CITY: Hackensack

; STATE: New Jersey

; COUNTRY: USA

; ZIP: 07601

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/849.303

; FILING DATE: 21-MAY-1997

; CLASSIFICATION: 514

; ATTORNEY/AGENT INFORMATION:

; NAME: Jackson Esq., David A.

; REGISTRATION NUMBER: 26,742

; REFERENCE/DOCKET NUMBER: 1321-1-003

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 201-487-5800

; TELEFAX: 201-343-1684

; TELEX: 133521

; INFORMATION FOR SEQ ID NO: 16:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 112 amino acids

; TYPE: amino acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: protein

; HYPOTHETICAL: NO

US-08-849-303-16

Query Match 20.6%; Score 136; DB 8; Length 112;
Best Local Similarity 30.6%; Pred. No. 5.1e-07;
Matches 34; Conservative 22; Mismatches 31; Indels 24; Gaps 5;

QY 22 NSPELEETLTHTITKLAENNATFYFKIDNVKKARVQVWAGKYFIDFVARETTCSEKN 81

Db 12 NEEGVQEALSPAVSEBFNKRSDAYQSRVVRVRARQVVGMYFLDVELGRITCTK--S 69

QY 82 EELTESC-----ETKLGQSLDCNAEYVVPWEKKIYPTVN-----CQ 119
Db 70 QANLDCPFHNPQHLKREKL-----CSFQYVYVVPWN-----TINLVKFSQ 111

Search completed: September 24, 2004, 14:13:03
Job time : 49.296 secs

